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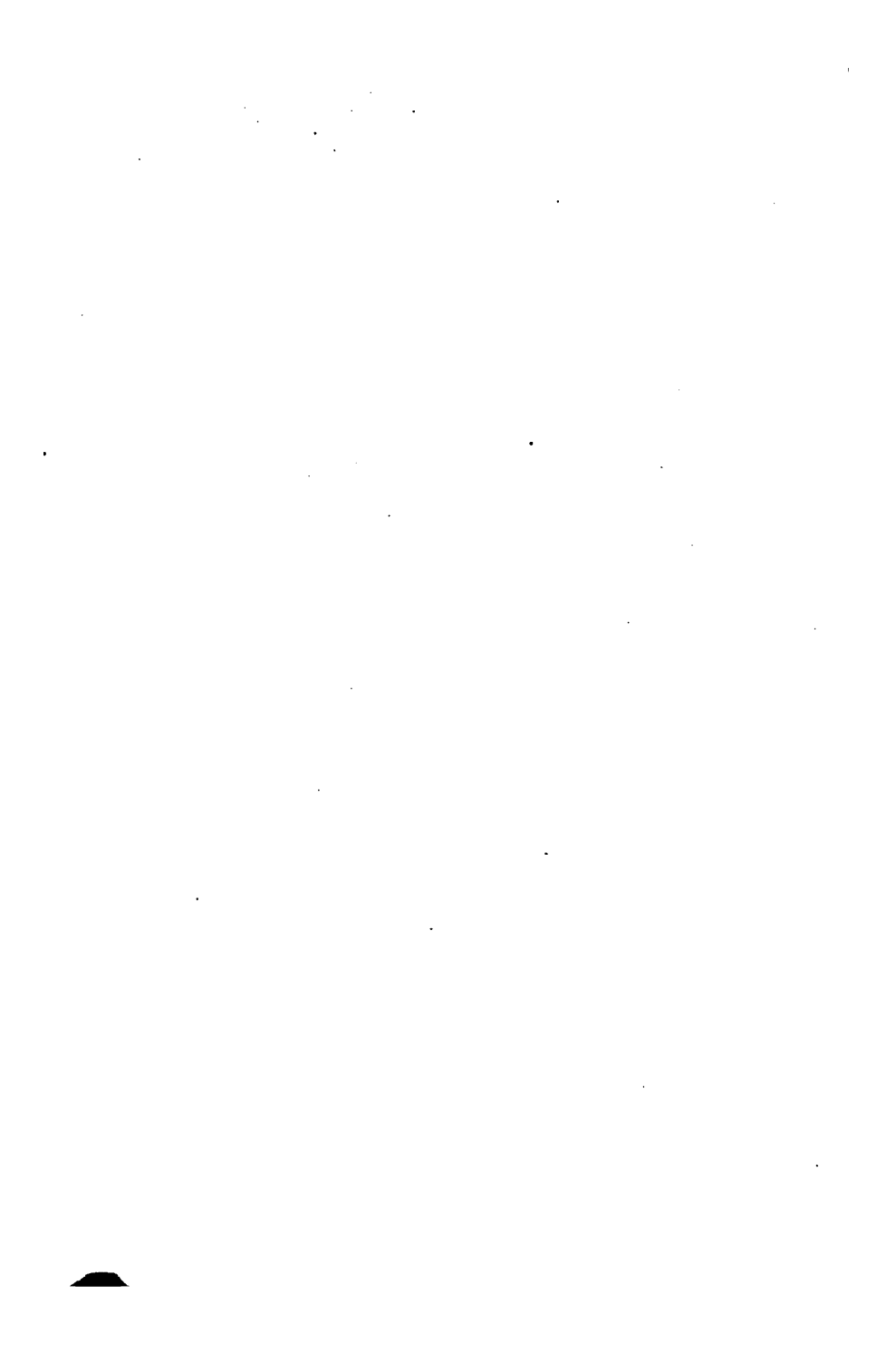
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CHEMISTRY:
GENERAL, MEDICAL, AND PHARMACEUTICAL;
INCLUDING
THE CHEMISTRY OF THE BRITISH PHARMACOPŒIA.

A MANUAL
ON THE GENERAL PRINCIPLES OF THE SCIENCE, AND THEIR
APPLICATIONS IN MEDICINE AND PHARMACY.

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193. f. 79.

"But the greatest error of all is, mistaking the ultimate end of knowledge; for some men covet knowledge out of a natural curiosity and inquisitive temper; some to entertain the mind with variety and delight; some for ornament and reputation; some for victory and contention; many for lucre and a livelihood; and but few for employing the Divine gift of reason to the use and benefit of mankind. Thus some appear to seek in knowledge a couch for a searching spirit; others, a walk for a wandering mind; others, a tower of state; others, a fort, or commanding ground; and others, a shop for profit or sale, instead of a storehouse for the glory of the Creator and the endowment of human life."—**Lord Bacon.**

P R E F A C E .

THE short title on the back of a book, and even the words on the title-page, are generally, and even necessarily, imperfect descriptions of the contents, and hence not unfrequently induce at the outset misconceptions in the minds of readers. The author of "Chemistry : General, Medical, and Pharmaceutical" would at once state, therefore, that his sole aim is to teach the general truths of chemistry to medical and pharmaceutical pupils. So far as laws and principles are concerned, the book is a work on General Chemistry ; but inasmuch as those laws and principles are elucidated and illustrated by that large portion of chemistry which is directly interesting to medical practitioners and pharmacists, the book may be said to be a work on Medical Chemistry and on Pharmaceutical Chemistry. Only in this conventional sense would the author speak of Medical and Pharmaceutical Chemistry ; for the truths of chemistry are the same for all students—crystalline verities which cannot be expanded or compressed to suit any class of workers. The leading principles of the science, however, can as easily be illustrated by or deduced from those facts which have interest as from those which have little or no special interest to the followers of medicine and pharmacy. The grand and simple leading truths or laws of chemistry, the lesser truths or principles, and nearly all the interest-

ing relationships of elements and compounds, in a word the *science* of chemistry, can be taught to medical and pharmaceutical students with little other aid than that afforded by the materials which lie in rich abundance all around these workers. Such a mode of teaching "the general principles of the science and their applications in medicine and pharmacy" is adopted in this volume. It is a mode which greatly increases the usefulness of the science to the class addressed, while it in no way diminishes the value of chemistry as an instrument of mental culture—an instrument which sharpens and expands the powers of observation, which enlarges and strengthens memory and imagination, which gives point to the perceptive faculties, and which develops and elaborates the powers of thought and of reason.

This manual is intended, then, as a systematic exponent of the general truths of chemistry, but is written solely for the pupils, assistants and principals engaged in medicine and pharmacy. It will be found equally useful as a reading-book for gentlemen having no opportunities of attending lectures or performing experiments, or, on the other hand, as a text-book for college pupils; while its comprehensive Index, containing nearly seven thousand references, will fit the work for after-consultation in the course of business or professional practice.

From other chemical text-books it differs in three particulars:—first, in the exclusion of matter relating to compounds which at present are only of interest to the scientific chemist; secondly, in containing more or less of the chemistry of every substance recognised officially, or in general practice, as a remedial agent; thirdly, in the paragraphs being so cast that the volume may be used as a guide in studying the science experimentally.

The order of subjects is that which, in the author's opinion, best meets the requirements of medical and pharmaceutical students in Great Britain, Ireland, America, and the English Colonies. Introductory pages are devoted to a few leading properties of the elements. A review of the facts thus unfolded affords opportunity for stating the views of philosophers respecting the manner in which these elements influence each other as components of terrestrial matter. The consideration in detail of the relations of the elementary and compound radicals follows, synthetical and analytical bearings being pointed out, and attention frequently directed to connecting or underlying truths or general principles. The chemistry of substances naturally associated in vegetables and animals is next considered. Practical toxicology and the chemical as well as microscopical characters of morbid urine, urinary sediments, and calculi are then given. The concluding sections form a laboratory-guide to the chemical and physical study of quantitative analysis. In the Appendix is a long table of tests for impurities in medicinal preparations—also a short one of the saturating-powers of acids and alkalies, designed for use in prescribing and dispensing.

In the course of the treatment outlined in the preceding paragraph, it will be observed that the whole of the elements are first noticed very shortly, to give the pupil a general view of his course of study, and afterwards at length and thoroughly; that the chemistry of the common metallic radicals precedes that of the rarer, and that the sections on the acidulous radicals are similarly divided; while the basylous radicals are arranged according to analytical relations, the common acidulous according to exchangeable value or quantivalence, and the rarer

acidulous radicals alphabetically. By this plan the more important facts and principles are repeatedly brought under consideration, the points of view, however, differing according as interest is concentrated on physical, synthetical, analytical, or quantitative properties. This arrangement of matter was adopted, also, partly from the belief that the separate and general truths of chemistry never enter the mind in the order of any scientific classification at present possible. In the current state of chemical knowledge, consistency in the methodical arrangement even of elements can only be carried out in one direction, and is necessarily accompanied by inconsistencies in other directions—a result most perplexing to learners, and hence totally subversive of the chief advantage of classification. For this reason the writer has preferred to lead up to, rather than follow scientific classification—has allowed analogies and affinities to suggest, rather than be suggested by, classification. Among the acidulous radicals, especially, any known system of classification would have given undue prominence to one set of relations, and undeserved obscurity to others. Then, by separating more important from less important matter, instruction is adapted to the wants of gentlemen whose opportunities of studying chemistry vary greatly, and are unavoidably insufficient to enable them to gain a knowledge of the detail of the science. One great advantage of the mode of treatment is that difficulties of nomenclature, notation, chemical constitution, and even those arising from conventionality of language, are explained as they arise, instead of being massed under the head of "Introductory Chapters," "Preliminary Considerations," or "General Remarks," which are not unfrequently too difficult to be understood by a beginner, too voluminous

to be remembered except by the aid of subsequent lessons, and are consequently the cause of much trouble and confusion. This plan has also admitted of greater prominence being given to "The General Principles of Chemical Philosophy," the only section to which the student is asked frequently to return until he finds himself naturally employing those principles in the interpretation of the phenomena obtained by experiment. An elementary knowledge of the subjects of Gravitation, Heat, Light, Sound, Electricity, and Magnetism cannot be too strongly recommended to the student of chemistry. The first portion of this manual would have been devoted to an exposition of these branches of physics, so far as they bear on chemistry, did not the many special books on physics render such a course unnecessary. Quantitative chemical analysis frequently involving determinations of temperature, specific gravity, and atmospheric pressure, a few paragraphs on these subjects are made introductory to the sections on quantitative operations.

The theories that matter consists of molecules, and that molecules consist of atoms, are freely adopted in this book, the author believing that, in the present state of knowledge and education, philosophic conceptions regarding chemistry can only be taught to medical, pharmaceutical, and the great majority of general students by some such objective aid.

The chemical notation of the work is in accordance with modern theories. Equations illustrative of pharmacopœial processes have a name attached to each formula.

Chemical nomenclature has been modernized to the extent of defining the alkali-metal salts and the earthy compounds as those of potassium, sodium, ammonium, barium, calcium, magnesium, and aluminium, instead of

potash, soda, ammonia, baryta, lime, magnesia, and alumina. The author confidently believes that this change, founded on views now adopted by all prominent writers on chemistry, will be accepted and become popular in medicine and pharmacy. It is a step in the direction of simplicity and consistency, and involves far less hypothesis than is contained in the old system. The name nitrate of potash, for example, was based on the pure assumption that nitre contained oxide of potassium or potash and nitric anhydride, then erroneously termed nitric acid. By the modern name nitrate of potassium, all that is intended to be conveyed is that nitre contains the element common to all potassium compounds, and the group of elements common to all nitrates. Under the old method, students always experienced difficulty in distinguishing salts of the metal from salts of its oxide—salts of potassium, for instance, from salts of potash; under the new view no such difficulty arises. Names such as potassium nitrate or potassic nitrate are also consistent with modern views, but for general adoption are too unlike the original. The contractions in Latin for names like “nitrate of potassium” are identical with the contractions for names resembling “nitrate of potash;” an accidental circumstance that will much facilitate the general introduction of the former names among medical practitioners and pharmacists, and a practical advantage that must determine the choice over the other chemically equivalent names just mentioned. The author ventures to express some gratification that his use and advocacy of this system since the first edition of this manual was published in 1867 has resulted in its adoption, in 1873, in the “Pharmacopœia of the United States,” and in the recommendation, from all medical, chemical and pharma-

ceutical authorities in Great Britain, of its adoption in the next "British Pharmacopœia." Pharmacy in these two countries will thus sooner or later, in the important matter of chemical nomenclature, be in accord with the current state of chemical science.

The Metric System of Weights and Measures (that which, doubtless, is destined to supersede all others) is alone used in the sections on Quantitative Analysis. In other parts of the manual avoirdupois weights and imperial measures are employed.

It is hoped that the numerous etymological references scattered throughout the following pages will be found useful. Words in Greek have been rendered in English characters, letter for letter. The word "official" is used throughout for things recognised officially by the compilers of Pharmacopœias; "officinal" in its original application to the officina or shop.

Students are strongly recommended to test their progress by frequent examination. To this end appropriate questions are appended to each subject.

The author's ideal of a manual of chemistry for medical and pharmaceutical students is one in which not only the *science* of chemistry is taught, but in which the chemistry of every substance having interest for the followers of medicine and pharmacy is noticed at more or less length in proportion to its importance, and at least its position in relation to the leading principles of chemistry set forth with all attainable exactness. The extent to which he has realized this ideal he leaves to others to decide. Such a work will doubtless in certain parts partake of the character of a dictionary; but this is by no means a fault, especially if a good index be appended; for the points of contact between pure and applied chemistry are

thus multiplied, and abundant outlets supplied, by which a lover of the science may pass into other chemical domains by aid of other guides, or even into the regions of original research. Among the rarer alkaloids, bitter bodies, glucosides, salts of organic radicals, solid fats, fixed oils, volatile oils, resins, oleo-resins, gum-resins, balsams, and colouring-matters mentioned in this volume will be found many such points whence the ardent student may start for the well-known, the obscure, or the untrodden paths of scientific chemistry.

Within thirteen years a demand has arisen for nine large editions of this manual. The First, in 1867, was intended as a handbook of practical chemistry only; but the notes and remarks made respecting most of the experiments were found to be so useful by students that this portion of the volume was in the Second Edition (1869) sufficiently extended to render the book more fairly complete in itself. In response to a call from professional friends in the United States in 1870, the work was revised by the author for the followers of medicine and pharmacy in America, the chemistry of the Preparations and Materia Medica of the United States Pharmacopœia being introduced, and such other adaptations included as to form a Third Edition. A Fourth was presented to English workers in 1872, and, founded on the fourth, a Fifth Edition for American students in 1873. A very large Sixth Edition was published in England in 1875, and, in America, a Seventh in 1876 and an Eighth in 1879.

The present, Ninth, edition contains such alterations

and additions as seemed necessary for the demonstration of the latest developments of chemical principles and the latest applications of chemistry to pharmacy.

About five hundred new references have been added to the Index, raising the total number to nearly seven thousand.

As the book has always included the chemistry of all articles officially recognised in the British Pharmacopœia, it has necessarily included nearly all the chemistry of the Pharmacopœia of India: into the sixth edition, however, was inserted some notice of each of forty substances official in the Indian, but not in the British Pharmacopœia. The work now includes the whole of the chemistry both of the British Pharmacopœia and of the Pharmacopœia of India.

Engravings of most of the important pieces of apparatus employed in studying chemistry experimentally have been inserted. The sixty or seventy blocks were cut by G. Pearson, Esq.; the twenty lithographs are by Tuffen West, Esq.

I am indebted to my assistants, Dr. Senier and Mr. Dunstan, for much valuable help in preparing the edition for the press. Dr. De Vrij has kindly revised the sections relating to alkaloids of the cinchonas.

17, BLOOMSBURY SQUARE, LONDON,
October 1st, 1880.

ADVICE TO STUDENTS.

It is unnecessary to advise you to avoid studying *merely* by way of "preparation for examination." You will not so mistake the means for the end. You are studying to fit yourself for your position in the world. Work diligently, study thoughtfully and deliberately, above all be thorough; otherwise your knowledge will be transient, and will be unaccompanied by that enlightenment of the understanding, that mental training, mental discipline, and general elevation of the intellect which constitute, in a word, education. When you are thus educated, you will with ease and pleasure pass any examination in the knowledge you have thus acquired.

All authorities on education, whether statesmen, teachers, or examiners, regard "Examination," even by the most highly skilled "Board," with ample time at its disposal and a wide area from which to select questions, as but a partial test of knowledge, and an imperfect test of education. It is the best, however, that has been devised, and is especially useful when, following instead of leading education, it is restricted to the subjects of a well-defined, earnestly followed, compulsory public curriculum of study—a curriculum defined and directed by a competent representative body, wisely administered by properly qualified teachers, and earnestly followed by pupils possessing sound preliminary training.

Students! as men of honour, take care that any inefficiencies inseparable from "examination" are abundantly compensated by the extent and precision of your knowledge, and by the soundness and thoroughness of your whole education.

APPARATUS FOR EXPERIMENTS IN ANALYSIS.

List of Apparatus suitable for the three months' course of practical chemistry in the summer session of medical schools, or for any similar series of lessons—including the preparation of elementary gases, analytical reactions of common metals and acidulous radicals, analysis of single salts, chemical toxicology, and the examination of urine, urinary sediments, and calculi :—

One dozen test-tubes.	A 2-inch and a 3-inch evap. basin.
Test-tube stand.	Two porcelain crucibles.
Test-tube cleaning-brush.	Blowpipe.
A few pieces of glass tubing, 8 to 16 in. long, with a few inches of india-rubber tubing to fit.	Crucible tongs.
Small flask.	Round file.
Two small beakers.	Triangular file.
Two small funnels.	Small retort-stand.
Two watch-glasses.	Sand-tray.
Two or three glass rods.	Wire triangles.
Wash-bottle.	Platinum wire and foil.
Small pestle and mortar.	Test-papers.
A 2-pint earthenware basin.	Filter-paper.
	Towel.
	Two dozen corks.

(This set, packed in a case, can be obtained of any chemical-apparatus maker for 12s. 6d. to 15s.)

APPARATUS FOR EXPERIMENTS IN SYNTHESIS AND ANALYSIS.

A larger set, suitable for the performance of most of the synthetical as well as analytical experiments described in this manual :—

A set of evaporating-basins, of the following sizes :—	One pair of 8-inch brass crucible-tongs.
One 8½-inch. One 4-inch.	Two soup-plates.
One 7½-inch. Two 3-inch.	One flat-plate.
One 6½-inch.	Two spatula knives.
One retort-stand and three rings.	One pair of scissors.
Two test-glasses.	One round file.
One half-pint flask.	One triangular file.
Half a quire of filter-paper.	Half a pound of glass rod.
Two porcelain crucibles.	Half a pound of glass-tubing. [ing.
One measure-glass, 5 oz.	One foot of small india-rubber tub.
Blowpipe, 8-inch, Black's.	Three dozen corks of various sizes.
Two glass funnels.	Platinum wire and foil.
One doz. test-tubes (German glass).	Test-papers.
One test-tube brush.	A nest of three beakers

(This set, packed in a case, can be obtained of any chemical apparatus maker for 25s. to 30s.)

A sponge, towels, and note-book may be included.

FURNITURE OF A LABORATORY.

The following apparatus should be ready to the hand of students following an extended course of practical chemistry, in a room set apart for the purpose:—

A bench or table and stool.	Test-tube rack, two dozen holes.
Water-supply and waste-pipe.	Iron stand or cylinder for supporting large dishes.
A cupboard attached to a chimney with an outward draught.	Iron adaptors for fitting dishes to cylinder.
A furnace fed with coke; tongs, hot-plate or sand-bath, etc.	Pestle and mortar, 5 or 6 inches.
A waste-box.	One 6-inch funnel.
Shelves for chemicals and other materials in jars or bottles.	Brown pan, 1 or 2-gallon.
Gas-supply and lamp with flexible tube (or a spirit-lamp and spirit).	White jug, 1-gallon.
	Water-bottle, quart.
	Twenty-eight test-bottles, 6 oz.

Other articles, such as flasks, retorts, receivers, condensers, large evaporating-dishes, may be obtained as wanted. In Quantitative Analysis the apparatus described in the sections on that subject will be required.

REAGENTS.

Certain chemicals are used so frequently in analytical processes that it is desirable to have small quantities placed in bottles in front of the operator. As these reagents or "tests" are generally employed in a state of solution, nearly all the solid salts may at once be dissolved (in distilled water). The bottles employed should be well stoppered, and of 5 or 6 ounces capacity. German-glass bottles of this size may be had for about four shillings per dozen. The bottles should not be more than about three-quarters full; single drops, if required, can then be poured out with ease and precision. The following list of test solutions is recommended; directions for methods of preparing those not readily purchasable will be found by referring to the Index:—

Sulphuric Acid, strong.	Sol. of Potash, 5 per cent. or B.P.
Nitric Acid, strong.	" Soda, 5 to 15 per cent.
Hydrochloric Acid, strong.	" Ammon., 10 per cent. or B.P.
Acetic Acid, strong.	Lime Water, saturated.

The next nine may contain about 10 per cent. of solid salt:—

Carbonate of Ammonium, with a little solution of Ammonia added.	Sulphhydrate of Ammonium.
Chloride of Ammonium.	Chloride of Barium.
Phosphate or Arseniate of Ammonium.	Chloride of Calcium.
	Phosphate of Sodium.
	Neutral Chromate.

The succeeding seven may have a strength of about 5 per cent. :-

Ferrocyanide of Potassium.	Perchloride of Iron.
Ferridecyanide of Potassium.	Nitrate of Silver.
Iodide of Potassium.	Perchloride of Platinum.
Oxalate of Ammonium.	

LISTS OF CHEMICALS.

List of chemicals necessary for the practical study of the non-metallic elements mentioned on pp. 1. to 22. The quantities are sufficient for several experiments.

Chlorate of Potassium	1 oz.	Phosphorus	$\frac{1}{2}$ oz.
Black Oxide of Manganese	1 oz.	Hydrochloric Acid	1 oz.
Zinc	1 oz.	Sulphur	$\frac{1}{2}$ oz.
Oil of Vitriol	2 oz.	Iodine	$\frac{1}{4}$ oz.

List of chemicals necessary for the *analytical* study of the metallic and acidulous radicals (pages 80 to 444). The quantities will depend on the frequency with which experiments are repeated or analysis performed; those mentioned are sufficient for one or two students. The articles are given in the order in which they will be required. The eight substances mentioned in the above list are included.

The set of test solutions described on the previous page.		Calomel	$\frac{1}{2}$ oz.
Carbonate of Potassium	1 oz.	Tin	1 oz.
Tartaric Acid	1 oz.	Bicarbonate of Potassium	1 oz.
Litmus	$\frac{1}{2}$ oz.	Acetate of Lead	1 oz.
Sulphate of Magnesium	1 oz.	Cyanide of Potassium	$\frac{1}{2}$ oz.
Sulphate of Zinc	1 oz.	Hyposulphite of Sodium	1 oz.
Alum	1 oz.	A Lithium Salt	10 grs.
Sulphide of Iron	1 lb.	Nitrate of Strontium	$\frac{1}{2}$ oz.
Oak-galls	1 oz.	Black Oxide of Manganese	$\frac{1}{2}$ lb.
Sulphocyanate of Potassium	$\frac{1}{2}$ oz.	Chloride of Manganese	$\frac{1}{2}$ oz.
White Arsenic	$\frac{1}{2}$ oz.	Chloride of Cobalt	50 grs.
Zinc	$\frac{1}{2}$ lb.	Nitrate of Nickel	$\frac{1}{2}$ oz.
Charcoal	$\frac{1}{2}$ lb.	Chloride of Chromium	$\frac{1}{2}$ oz.
Sulphate of Iron	1 oz.	Gold leaves	2 or 3
Copper Foil	1 oz.	Chloride of Cadmium	$\frac{1}{2}$ oz.
Sulphate of Copper	1 oz.	Nitrate of Bismuth	$\frac{1}{2}$ oz.
Tartar Emetic	$\frac{1}{2}$ oz.	Bromide of Potassium	$\frac{1}{2}$ oz.
Mercury	1 oz.	Starch	1 oz.
Corrosive Sublimate	$\frac{1}{2}$ oz.	Nitrate of Potassium	1 oz.
		Copper borings or turnings	1 oz.
		Indigo	$\frac{1}{2}$ oz.

Chlorate of Potassium	1 oz.	Borax	1 oz.
Iodine	$\frac{1}{2}$ oz.	Turmeric	1 oz.
Spirit of Wine	1 oz.	Benzoic Acid	.50 grs.
Sulphur	1 oz.	Fluor Spar	1 oz.
Acid Oxalate of Potassium	1 oz.	Tannic Acid	.50 grs.
Citric Acid	1 oz.	Gallie Acid	.50 grs.
Phosphorus	1 oz.	Pyrogallie Acid	.50 grs.

The quantities of materials required for the study of chemistry *synthetically* will necessarily vary with the desires and tastes of the operator, or according to the number and requirements of students working together.

ERRATA.

PAGE.	LINE.	
155	30	for K_3FeO_4 read K_2FeO_4 .
177	6	for HN_4HS read NH_4HS .
228	15	dele "of the."

CONTENTS.

	PAGE
PREFACE	iii
ADVICE TO STUDENTS	xiii
APPARATUS	xv
REAGENTS	xvi
LISTS OF CHEMICALS	xvii
INTRODUCTION	1
GENERAL PROPERTIES OF THE NON-METALLIC ELEMENTS	3
SYMBOLS AND DERIVATION OF NAMES OF ELEMENTS	23
THE GENERAL PRINCIPLES OF CHEMICAL PHILOSOPHY	29
COMMON METALLIC ELEMENTS, THEIR OFFICIAL PREPARATIONS AND TESTS :—	
SALTS OF POTASSIUM, SODIUM, AMMONIUM, BARIUM, CALCIUM, MAGNESIUM, ZINC, ALUMINIUM, IRON, ARSENICUM, ANTIMONY, COPPER, MERCURY, LEAD, SILVER	59
ANALYTICAL CHARTS FOR ORDINARY METALS	252
RAREER METALLIC ELEMENTS, THEIR OFFICIAL PREPARATIONS AND TESTS :—	
SALTS OF LITHIUM, STRONTIUM, MANGANESE, COBALT, NICKEL, CHROMIUM, TIN, GOLD, PLATINUM, CADMIUM, BISMUTH	257
ANALYTICAL CHARTS FOR ALL METALS	296
COMMON ACIDULOUS RADICALS, OFFICIAL ACIDS, AND TESTS :—	
CHLORIDES, BROMIDES, IODIDES, CYANIDES, NITRATES, CHLORATES, ACETATES, SULPHIDES, SULPHITES, SULPHATES, CARBONATES, OXALATES, TARTRATES, CITRATES, PHOSPHATES, BORATES	302
SALTS OF RAREER ACIDULOUS RADICALS :—	
BENZOATES, CYANATES, FORMATES, HIPPURATES, FERROCYANIDES, FERRIDCYANIDES, FLUORIDES, HYPOPHOSPHITES,	

	PAGE
HYPOSULPHITES, LACTATES, MALATES, MECONATES, METAPHOSPHATES, NITRITES, PHOSPHITES, PYROPHOSPHATES, SILICATES, SULPHOCYANATES, TANNATES, GALLATES, URATES, VALERIANATES	392
ANALYTICAL CHART FOR ACIDULOUS RADICALS	431
SYSTEMATIC ANALYSIS	433
ALKALOIDS, AMYLACEOUS, AND SACCHARINE SUBSTANCES, GLUCOSIDES, ALCOHOL AND ALLIED BODIES, ALBUMENOID AND GELATINOGENOUS SUBSTANCES, PEPSIN, FATTY BODIES, RESINOID SUBSTANCES, COLOURING-MATTERS	445
TOXICOLOGY	577
EXAMINATION OF MORBID URINE AND CALCULI	590
OFFICIAL GALENICAL PREPARATIONS	605
OFFICIAL CHEMICAL PREPARATIONS	607
QUANTITATIVE ANALYSIS :—	
INTRODUCTORY REMARKS	608
MEASUREMENT OF ATMOSPHERIC PRESSURE	610
MEASUREMENT OF TEMPERATURE	611
ESTIMATION OF WEIGHT	618
WEIGHTS AND MEASURES	619
SPECIFIC GRAVITY	627
CORRECTION OF THE VOLUME OF GASES FOR PRESSURE AND TEMPERATURE	635
VOLUMETRIC QUANTITATIVE ANALYSIS	643
GRAVIMETRIC QUANTITATIVE ANALYSIS	673
DIALYSIS	729
APPENDIX :—	
TABLE OF OFFICIAL TESTS FOR IMPURITIES IN PREPARATIONS OF THE BRITISH PHARMACOPEIA	735
SATURATION TABLES	743
THE ELEMENTS, THEIR SYMBOLS, AND ATOMIC WEIGHTS	744
INDEX	747

CHEMISTRY:

GENERAL, MEDICAL, AND PHARMACEUTICAL.

INTRODUCTION.*

THE infinite variety of solid, liquid, and gaseous substances of which our earth and atmosphere are composed may be resolved, with more or less difficulty, into distinct forms of matter appropriately termed Elements, for by no known means can they be further decomposed. Sixty-seven Elements have been proved to exist. A few (such as gold) occur naturally in the uncombined state; but the greater number are combined in so subtle a manner as to conceal them from ordinary methods of observation. Thus none of the common properties of water indicate that it is composed of two elements, both gases, but differing much from each other: nor can the senses of sight, touch, and taste, or other common means of examination, detect in their concealment the three elements of which sugar is composed. The art by which these and all other compound substances are resolved into their elements is termed Chemistry, a name derived possibly from the Arabic word *kamai*,

* Students using this book as a guide in following chemistry practically, should read the first four pages, and then commence work by preparing oxygen. All students should read the Preface.

to conceal.* The *art* of chemistry also includes the construction of compounds from elements, and the conversion of substances of one character into those of another. The general principles or leading truths relating to the elements, to the manner in which they severally combine, and to the properties of the compound substances formed by their union, constitute the *science* of chemistry.†

From these few words concerning the nature of the art

* The idea that common metals contained valuable metals concealed within them was the one seed from which mainly sprung chemical knowledge. The men who endeavoured to find the secret of such concealment were appropriately termed *alchemists*, and their efforts spoken of as *alchemy* (*al kimia*, from *kamai*, to conceal). Their persistent labours, generation after generation, were unsuccessful so far as the transmutation of baser metals into gold was concerned, yet were invaluable to posterity; for new substances were discovered and truths of nature unveiled: from these discoveries multiplication of discoveries resulted, and thus grew the still growing branch of knowledge called Chemistry.

† Persons who practise the art and science of Chemistry are known as Chemists. Some two hundred or more years ago, and before chemistry was a science, the only "chemists" were the makers and vendors of chemicals, then only used as medicines. They were the successors of the Alchemists. In Great Britain these chemists and the herb-dealers, otherwise drug-grocers, otherwise drug-gists, gradually associated to form the "Chemist and Druggist." Between the "Chemist and Druggist" and the Physician there existed the Apothecary—the putter together of medicines, or compounder of physician's prescriptions. The Apothecary has since become a medical practitioner, prescriptions being now "made up" by the Chemist and Druggist. The latter in Great Britain, since the year 1868, is allowed the title of *Chemist and Druggist*, his higher title being *Pharmaceutical Chemist*; these respective designations he legally assumes on passing the Minor and Major Examinations conducted by the Pharmaceutical Society of Great Britain, in accordance with the provisions of the Pharmacy Acts of 1852 and 1868. The whole class is often spoken of as that of *Pharmacists* or *Pharmaceutists*, terms also used in the United States. Other classes of Chemists are the Analytical Chemists, who give special attention to analysis; Manufacturing Chemists, who restrict their labours to the preparation of chemicals; while others devote a portion of their knowledge and energies to Chemical Education or to Chemical Research, or are appealed to as Consulting Chemists by the persons, firms, corporations, or governments needing chemical advice respecting industrial processes, hygienic matters, etc. The callings of the Consulting and Analytical Chemist are generally united, and the professional gentlemen who follow these conjoint avocations also not unfrequently occupy professorial or other tutorial positions, sometimes adding to these labours more or less work at original chemical research. In England, Scotland, and Ireland, nearly all the leading professional chemists are Fellows of the Institute of Chemistry of Great Britain and Ireland.

and science of chemistry, it will be seen that in most of the occupations that engage the attention of man chemistry plays an important part—in few more so than in the practice of Therapeutics * and of Pharmacy.†

Air, water, food, drugs, and chemicals, in short, all material substances are composed, as stated, of Elements. An intimate knowledge of the properties of the more important Elements, and of the various substances they form by combining with each other, a knowledge of the power or force (the chemical force or chemical affinity) by which the elements contained in the compounds are held together, and an application of such knowledge to Pharmacy and Medicine, must be the objects sought to be attained by the learner for whom this book has been especially written.

The Elements.—Of the sixty-seven known elements, thirty-nine are of medical or pharmaceutical interest; of these, about two-thirds are metals, and one-third non-metals; the remainder are so seldom met with in nature as to have received no practical application, either in medicine, art, or manufacture. Before intimately studying the elements, it is desirable to acquire some general notions concerning them: such a procedure will also serve to introduce the practical student to his apparatus, and make him better acquainted with the various methods of manipulation.‡

* Therapeutics (θεραπευτικός, *therapeutikos*, from θεραπεύω, *therapeuō*, to nurse, serve or cure) is that branch of medicine which treats of the application of remedies for diseases; it includes dietetics. The therapist also takes cognizance of hygiene—that department of medicine which respects the preservation of health.

† Pharmacy (from φάρμακον, *pharmakon*, a drug) is the generic name for the operations of preparing or compounding medicines, whether performed by the Medical Practitioner or by the Chemist and Druggist. It is also sometimes applied, like the corresponding term Surgery, to the apartment in which the operations are conducted.

‡ This allusion to apparatus need not discourage the youngest pupil. With the aid of a few phials, wine-glasses, or other similar vessels, always at hand, he may, by studying the following pages, learn the chemical reactions which are constantly occurring in the course of making up medicine, understand the processes by which medicinal preparation are

Metallic Elements.—With regard to the metallic elements, it may safely be assumed that the reader has sufficient knowledge for present purposes; but little, therefore, need now be said respecting them. He has an idea of the appearance, relative weight, hardness, etc., of such metals as gold, silver, copper, lead, tin, zinc, and iron. If he has not a similar knowledge of mercury, antimony, arsenicum, platinum, nickel, aluminium, magnesium, potassium, and sodium, he should embrace the earliest opportunity of seeing and handling specimens of each of these metals.

*Non-Metallic Elements.**—With regard to the non-metallic elements, it is here supposed that the student has no general knowledge. He should commence his studies therefore by a series of operations as follows, on eight out of their number.

OXYGEN.

Preparation.—As oxygen is the most abundant element in nature, forming, though in a combined state, about one-half of the whole weight of our globe, it may safely be assumed that this element can readily be obtained in the free condition in a state of purity. In fact, the air itself contains about one-fifth of its bulk of oxygen, though from the air it cannot be separated, at least not easily and readily, for experimental purposes. It is preferable to apply heat—that force which will often be noticed as antagonistic, so to speak, to chemical union; heat generally separating particles of matter further from each other, while chemical attraction tends to bind them closer together—it is better to heat certain compounds containing oxygen; the latter is then evolved in its normal, natural condition of gas. Several substances, when heated, yield oxygen; but, for convenience and economy, the crystalline body known as chlorate of potassium is best fitted for the experiment. The

manufactured, and detect adulterations, impurities, or faults of manufacture. Among the substances used in medicine will be found nearly all the chemicals required. If, in addition, a dozen test-tubes and a few feet of glass tubing be procured, many of the experiments described may be performed. For full lists of apparatus, see the Introductory pages.

* These bodies are sometimes termed *metalloids* (from *μέταλλον*, *metallon*, a metal, and *εἶδος*, *eidos*, likeness); but the name is not appropriate, for the non-metallic elements have no likeness to metals.

size and form of the vessel in which to heat it will mainly depend on the quantity required; but for the purposes of the student the best is a *test-tube*, an instrument in constant requisition in studying practical chemistry. It is simply a thin tube of glass, a few inches in length, and half or three-quarters of an inch in diameter, closed by fusion at one end. It is made of thin glass, in order that it may be rapidly heated or cooled without risk of fracture. (See figs. 3 and 4.)

Outline of the Process.—Heat chlorate of potassium (say, as much as will lie on a shilling) in a test-tube, by means of a spirit- or gas-flame; gaseous oxygen is quickly evolved. Before applying heat, however, provision should be made for collecting the gas.

Collection of Gases (see fig. 3).—Procure a piece of glass tubing about the thickness of a quill pen, and a foot or eighteen inches long, and fit it accurately to the test-tube by means of a cork. (Longer tubes may be neatly cut to any size by smartly drawing the edge of a triangular file across the glass at the required point, then clasping the tube, the scratch being between the hands, and pulling the portions asunder, force being exerted in a *slightly* curved direction, so as to open out the crack which the file has commenced.) The tube is fixed in the cork through a round hole made by the aid of a red-hot wire, or, better, by a rat-tail file, or, best of all, by one of a set of cork-borers—pieces of brass tubing sharpened at one end and having a flat head at the other. Setting aside the test-tube for a few minutes, proceed to bend the long piece of tubing to the most convenient shape for collecting the gas.

Fig. 1.

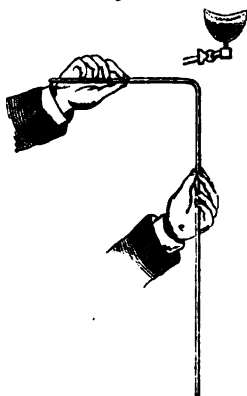


SOFTENING AND BENDING GLASS TUBES.

To Bend Glass Tubes.—Hold the part of the tube required to be bent in any gas- or spirit-flame (a fish-tail gas-jet answers

very well), constantly rotating it, so that about an inch of the glass becomes heated. It will soon be felt to soften, and will then, yielding to the *gentle* pressure of the fingers, assume any required angle. In the present case, the tube should be heated at about four inches from the extremity to which the cork is attached, and bent to an angle of about 90 degrees.

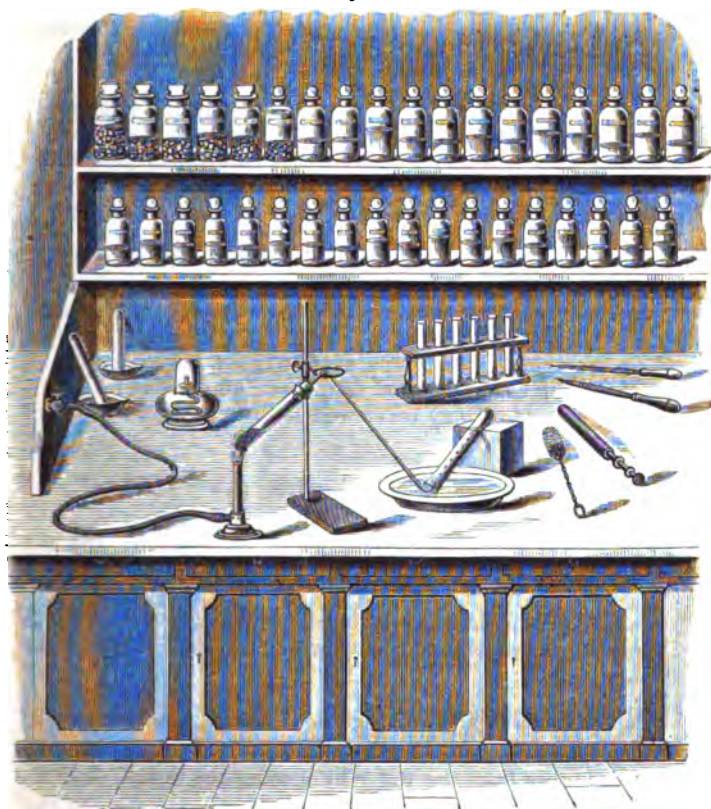
Fig. 2.



Source of Heat.—The source of heat for the test-tube may be the flame of an ordinary spirit lamp, or, still better where coal-gas is procurable, a mixture of the latter with air. Gas-lamps especially constructed to burn a mixture of coal-gas and air are sold by chemical apparatus manufacturers. (See figs. 3 and 7.)

Collection, etc. (continued).—Fit the cork and bent tube into the test-tube; the apparatus will then be ready for delivering gas at a convenient distance from the heated portion of the arrangement. To collect it, have ready three or four test-tubes (or small wide-mouthed bottles) filled with water, and inverted in a basin, or other similar vessel, also containing water, taking care to keep the mouths of the tubes a little below the surface. Now apply heat to the chlorate contained in the test-tube, and so arrange the open end of the bent tube under the water that the gas which presently issues may bubble into and gradually fill the inverted test-tubes. The first tubeful may be rejected, as it probably consists of little more than the air originally in the apparatus, and which has been displaced by the oxygen. That which comes afterwards will be pure oxygen.

As each tube or bottle becomes full, its mouth (still under the surface of the water) may be closed by a cork and set aside; or a little cup (such as a porcelain crucible or small gallipot) may be brought under the mouth, and the cup, with

Fig. 3.

This engraving represents the preparation, collection, and storage of small quantities of oxygen gas. A test-tube and bent glass tube, joined together by a perforated cork, are supported by the arm of an iron stand. (The apparatus might be held by the fingers.) The tube is heated by a gas-lamp. (The spirit-lamp shown at back might be used instead.) Gas evolved from the heated substance in the test-tube is displacing water from an inverted test-tube. Spare tubes in a test-tube rack are at hand, and tubes already filled are set aside till wanted. A nest of cork-borers, a round file, a triangular file, and a test-tube cleaning-brush are lying on the table or student's bench. Below are cupboards for apparatus, above are bottles containing testing liquids, etc.

the mouth of the tube in it, be lifted out of the water and placed close by till wanted, the water remaining in the cup effectually preventing the gas from escaping.

On the large scale, oxygen may be made in the same way, larger vessels (glass flasks or iron bottles) being employed. Less heat also will be necessary if the chlorate of potassium be previously mixed with very fine sand, or, still better, with about an equal weight of common black oxide of manganese.

Note on the Collection and Storage of Gases.—It may be as well to state that nearly all gases, whether for experimental or practical purposes, are collected and stored in a similar manner. Even coal-gas is generated at gas-works in iron retorts very much the shape of test-tubes, only they are as many feet long as a test-tube is inches; and the well-known gigantic gas-holders may be viewed as inverted iron test-tubes of great diameter.

Properties.—Oxygen is a colourless gas. Cailletet and Pictet have succeeded in liquefying it. Obviously it is not very soluble in water, or it could not be collected by the aid of that liquid.

Oxygen is soluble to a certain extent, however (about 3 volumes in 100, at common temperatures), or fishes could not breathe.

Other noticeable features are its want of taste and smell. Next, to show the relation of oxygen to combustion, remove one of the tubes from the water by placing the thumb over its mouth, and apply for a second a lighted wood match to the orifice; the gas will be found to be incombustible. Extinguish the flame of the match, and then quickly introduce the still incandescent carbonaceous extremity of the wood half-way down the test-tube; the wood will at once burst into flame, owing to the extreme violence with which oxygen supports combustion. These tests of the presence of oxygen may also be applied at the extremity of the delivery tube whilst the gas is being evolved. (It is desirable to retain two tubes of the gas for use in subsequent experiments; also one tube in

which only one-third of the water has been displaced by oxygen.)

Relation of Oxygen to Animal and Vegetable Life.—Not only the carbon at the end of a piece of charred wood, but any other substance that will burn in air (which, as will be seen presently, is diluted oxygen) will burn more brilliantly in pure oxygen. The warmth of the bodies of animals is kept up by the continuous burning of the tissues in the oxygen (of the air) drawn into the system through the lungs. The product of this combustion is exhaled into the air as a gaseous compound of carbon and oxygen termed carbonic acid gas, a gas which, in sunlight, is decomposed in the cells of plants with fixation of the carbon and liberation of the oxygen; hence the atmosphere is kept constant in composition.

Memorandum.—At present it is not advisable that the reader should trouble himself with the consideration of the chemical action which occurs either in the elimination of oxygen from its compounds, or in the separation of any of the following non-metallic elements from their combinations. It is to the properties of the elements themselves that he should restrict his attention. Working thus from simple to more complex facts, he will in due time find that the comprehension of such actions as occur in the preparation of these few elements will be easier than if he attempted their full study now.

HYDROGEN.

Preparation and Collection.—The element Hydrogen is also a gas,* and is obtainable from its commonest compound, water (of which one-ninth by weight is hydrogen), by the agency of hot zinc or iron, but more conveniently by the action of either of those metals on cold diluted sulphuric acid. The apparatus used for making oxygen may be employed for this experiment; but no lamp is required. Place several pieces

* Graham obtained alloys of hydrogen with palladium and other metals, compounds in which several hundred times its bulk of gas is retained by the metal in vacuo or even at a red heat. This was physical confirmation of the opinion long held by chemists, that hydrogen is a gaseous metal. Graham termed it *hydrogenium*, other chemists *hydrium*, and considered its relative weight in the solid state to be nearly three-fourths that of water. Cailletet and Pictet have since actually liquefied and solidified this element.

of thin zinc* in the generating tube (fig. 4), or in any common glass bottle (fig. 5) or flask, and cover them with water. The collecting tubes (these also may be wide-mouthed bottles) being ready, add strong sulphuric acid (oil of vitriol) to the zinc and water, in the proportion of about 1 volume of

Fig. 4.



Fig. 5.



PREPARATION OF HYDROGEN.

acid to 5 of water, and fit on the delivery-tube, or pour the acid down such a funnel-tube† as is shown in fig. 5; the hydrogen is at once evolved. Having rejected the first portions (or having waited until the air originally in the bottle may be considered to be all expelled), collect four or five tubes of the gas in the manner described under Oxygen.

This process is similar to that of the "British Pharmacopœia." In making larger quantities, bottles of appropriate size may be employed.

Other metals, notably potassium and sodium, liberate hy-

* The best form is *granulated zinc* (*Zincum Granulatum*, B. P.), made by heating scraps of common sheet zinc in a ladle over a fire, and, as soon as melted, pouring, in a slow stream, into a pail of water from a height of 8 or 10 feet. Each drop of zinc thus yields a thin little bell, which, for its weight, presents a large surface to the action of the acid liquid. If the zinc is allowed to become hotter than necessary, the little bells will not be formed.

† Funnel-tubes may be purchased of the apparatus-maker; or, if the pupil has access to a table blowpipe and the advantage of a tutor to direct his operations, they may be made by himself.

drogen the moment they come into contact with water; but the processes are not economical.

Properties.—Like oxygen, hydrogen gas is invisible, inodorous, and tasteless. If made with iron it has a strong smell; but this is due to impurities derived from the iron.

Apply a flame to the mouth of the delivery-tube (care being taken that the gas is coming off briskly—a guarantee that no air remains in the generating vessel); ignition of the hydrogen ensues, showing that, unlike oxygen, it is combustible.

Plunge a lighted match well into a tube (or wide-mouthed bottle) containing hydrogen; the gas is ignited, but the match becomes extinguished. This shows that hydrogen is not a supporter of combustion.

Hydrogen in burning unites with the oxygen of the air and forms water, which may be condensed on a cool glass or other surface. Prove this by holding a glass vessel a few inches above a hydrogen-flame. In burning the hydrogen contained in one of the tubes or bottles, the flame is best seen when the tube is held mouth upwards, and water poured in so as to expel the gas gradually.

If, instead of this gradual combination of the two elements oxygen and hydrogen, they be mixed together in bulk in the right proportions and then ignited, they will rapidly combine, and explosion will result. Prepare a mixture of this kind by filling up with hydrogen a test-tube from which one-third of the water has been expelled by oxygen. Remove the tube from the water, placing a finger over the mouth, and having a lighted match ready, apply the flame; a slight explosion ensues, owing to the instantaneous combination of the two elements, and the expansive force of the highly heated steam produced. If anything larger than a test-tube is employed in this experiment, it should be a soda-water bottle, or some such vessel equally strong.

These two gases thus unite at a temperature far higher than that of boiling water, two volumes of hydrogen and one

of oxygen yielding two volumes of gaseous water (true steam).

The noise of such explosions is caused by concussion between the suddenly expanded gaseous body and the air.

The force of the explosion, or, in other words, the expansive force of the suddenly heated and therefore suddenly expanded steam, is below that necessary to break the test-tube. Some force, however, is exerted; and hence the necessity of the precaution previously suggested of allowing all the air which may be in a hydrogen-apparatus to escape before proceeding with the experiments. If a flame be applied to the delivery-tube before all the air is expelled, the probable result will be ignition of the mixture of hydrogen and oxygen (of the air) and consequent explosion. But even in this case the generating vessel is not often fractured unless it be large and of thin glass, the ordinary effect being that the cork is blown out, and the delivery-tube broken on falling to the ground.

Hydrogen is an important constituent of all the substances used for producing artificial light, such as tallow, oil, and coal-gas. The explosive force of large quantities, such as a roomful, of coal-gas and air, though vastly below that of an equal weight of gunpowder, is well known to be sufficient at least to blow out that side of the room which offers least resistance.

The composition of water can be proved analytically as well as synthetically, a current of electricity decomposing it into its constituent gases, twice as much hydrogen as oxygen, by volume, being produced.

Combustion (from *comburo*, to burn).—The experiments with hydrogen and oxygen illustrate the true character of combustion. Whenever chemical combination is sufficiently intense to be accompanied by heat and light, the materials are said to undergo combustion. Combustion only occurs at the line of contact of the combining bodies; a jet of oxygen will burn in an atmosphere of hydrogen quite as easily as a jet of hydrogen in oxygen. A jet of air (diluted oxygen) will burn as readily in a jar of coal-gas as a jet of coal-gas burns in air; each is combustible, each supports the combustion of the other. Hence the terms *combustible* and *supporter of combustion* are purely conventional, and only applicable so long as the circumstances under which they are applied remain the same. In the case of substances burning

in air, the conditions are, practically, always the same; hence no confusion arises from regarding air as the great supporter of combustion, and bodies which burn in it as being combustible.

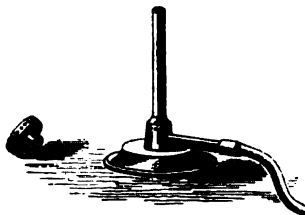
Structure of Flame.—A candle-flame (fig. 6) or oil-flame is a jet of gas intensely heated; the central portion is unburnt gas, the next envelope is formed of partially burnt and very dense gaseous and solid particles sufficiently highly heated to give light, and the outer cone of completely burnt gases. In the figure the sharpness of limit of these cones is purposely somewhat exaggerated. Air made, by any mechanical contrivance of burner, to mix with the interior of a flame at once burns up, or perhaps prevents the formation of dense gases, giving a hotter, but non-luminous jet. The air-gas lamps (fig. 7), or "Bunsen" gas-burners, commonly used in chemical laboratories are constructed on this principle: their flame has the additional advantage of not yielding a deposition of soot.

Fig. 6.



STRUCTURE OF FLAME.

Fig. 7.



"BUNSEN," OR AIR-GAS, BURNER.

In the air-gas lamp coal-gas escaping from a small orifice draws rather more than twice its volume of air (supplied through adjacent holes) into its column, and the mixture of gas and air passes upwards along a pipe. It only burns at the end, and not within the pipe, partly because the metal of the burner, by conducting heat away, cools the mixture below the temperature at which it can ignite; partly because the velocity with which the mixture flows out is greater than the rate at which such a mixture ignites; and partly because the

proportion of air to gas in the mixture is insufficient for thorough and perfect combustion, the external air contributing materially to the complete combustion of the jet of air-gas. The *Davy safety-lamp* acts on the principle first named: a wire-gauze cage surrounds an oil-flame; an inflammable mixture of gas (fire-damp) and air can pass through the gauze and catch fire and burn inside; but the flame cannot be communicated to the mixture outside, because the metal of the gauze and of the other parts cools down the gas below the temperature at which combustion can continue.

Properties (continued).—Gaseous hydrogen is the lightest substance known. It was formerly used for filling balloons, but was superseded by coal-gas. Coal-gas is not so light as hydrogen, but is cheaper and more easily obtained. The lightness of hydrogen may be rendered evident by the following experiment:—Fill two test-tubes with the gas, and hold one with its mouth downwards and the other with its mouth upwards. The hydrogen will have escaped from the latter in a few seconds, whereas the former will still contain the gas after the lapse of some minutes. This may be proved by applying a lighted match to the mouths of the respective tubes.

The relative weight or specific gravity of oxygen is sixteen times that of hydrogen. A vessel holding one grain of hydrogen will hold sixteen grains of oxygen. The relation of the weight of hydrogen to air is as 1 to 14.44 or as 0.0693 to 1.0. One grain of hydrogen by weight would measure about 27 fluid ounces, and, therefore, would about fill a common English wine-bottle. Such a bottle would, at common temperatures, hold about $14\frac{1}{2}$ grains of air, or 16 grains of oxygen.

Mem.—It is desirable to retain two tubes of hydrogen for use in subsequent experiments.

Diffusion of Gases.—Hydrogen cannot be kept in such vessels as the inverted test-tube; for, though much lighter than air, it *diffuses* downwards into the air, while the air, though much heavier, diffuses upwards into the hydrogen. This power of *diffusion* is characteristic of all gases, and proceeds according to a fixed rate, namely, “in inverse proportion to the square root of the specific gravity of the gas” (*Graham*). Thus hydrogen diffuses four times faster than oxygen. This

great and important property of diffusion strongly suggests that the particles of gases, at least, are always moving, never at rest; how otherwise *could* gases diffuse into each other as they do, notwithstanding the opposing influence of gravitation. Diffusion strongly supports this (Clausius's) kinetic (*κίνησις*, *kineō*, I move, or put in motion) theory of the physical condition of gases.

PHOSPHORUS.

Appearance and Source.—Phosphorus (*Phosphorus*, B.P.) is a solid element, in appearance and consistence resembling white wax; but it gradually becomes yellow by exposure to light. It is a characteristic constituent of bones, and is always prepared from that source by a process which will be subsequently described.

Caution.—Phosphorus, on account of its great affinity for oxygen, takes fire very readily in the air, and should therefore be kept under water. When wanted for use it must be cut under water. It is employed in tipping lucifers, though red or amorphous phosphorus (*vide* Index) is least objectionable for this purpose.

Experiment.—Dry a piece about one-fourth the size of a pea by quickly and carefully pressing it between the folds of porous (filter or blotting) paper; place it on a plate, and ignite by touching it with a piece of warm wire or wood. The product of combustion is a dense white suffocating smoke, which must be confined at once by placing an inverted tumbler, test-glass, or other similar vessel over the phosphorus. The fumes rapidly aggregate, and fall in white flakes on the plate. When this has taken place, and the phosphorus is no longer burning, moisten the powder with a drop or two of water, and observe that some of the water is converted into steam, an effect due to the intense affinity with which another portion of the water and the powder have combined.

The powder produced by the combustion of phosphorus is phosphoric anhydride; the combination of the latter with the elements of water produces a variety of phosphoric acid which dissolves in the water, forming, on standing, a dilute solution

of ordinary phosphoric acid. The Diluted Phosphoric Acid of the "British Pharmacopœia" is a somewhat similar solution, made, however, in a different way, and of a definite strength.

NITROGEN.

Source.—The chief source of this gaseous element is the atmosphere, nearly four-fifths of which consist of nitrogen (the remaining fifth being almost entirely oxygen).

Preparation.—Burn a piece of dried phosphorus, the size of a pea, in a confined portion of air. The oxygen is thus removed, and nitrogen alone remains. The readiest mode of performing this experiment is to fix a piece of earthenware (the lid of a small porcelain crucible answers very well) on a thin piece of cork, so that it may float in a dish of water (fig. 8). Place the phosphorus on the lid, ignite by a warm rod, and then invert a tumbler, or any glass vessel of about a half-pint capacity, over the burning phosphorus, so that the mouth of the glass may dip into the water. Let the arrangement rest for a short time for the fumes of phosphoric anhydride to subside and dissolve in the water, and then decant the gas into test-tubes in some such manner as that indicated in fig. 9, using a tub or other vessel of water of sufficient depth to admit of the glass containing the nitrogen to be turned on one side without air gaining access.

Fig. 8.



PREPARATION OF NITROGEN.

Fig. 9.



DECANTATION OF GASES.

Larger quantities of nitrogen are made in the same way. Other combustibles, such as sulphur or a candle, might be used to burn out the oxygen from a given quantity of air, but none answer so quickly and completely as phosphorus; added to which, the product of their combustion would not always be dissolved by water, but would remain with and contaminate the nitrogen.

Mem.—The statement concerning the composition of the air is roughly confirmed in preparing nitrogen, about one-fifth of the volume of the air originally in the glass vessel having disappeared, its place being occupied by water from the dish.

Properties.—Like oxygen and hydrogen, nitrogen gas is invisible, tasteless, and inodorous. By pressure, Cailletet and Pictet have succeeded in condensing it to a liquid. It is only slightly soluble in water. *Free* nitrogen is distinguished from all other gases by the absence of any characteristic or positive properties. Apply a flame to some contained in a tube; it will be found to be incombustible. Immerse a lighted match in the gas; the flame is extinguished, showing that nitrogen is a non-supporter of combustion.

The chief office of nitrogen in the air is to dilute the energetic oxygen, a mere *mechanical* mixture resulting.

Nitrogen is fourteen times as heavy as hydrogen.

The air is nearly fourteen and a half (14.44) times as heavy as hydrogen. Its average composition, including minor constituents (which will be referred to subsequently), is as follows:—

Composition of the Atmosphere.

	In 100 volumes.
Oxygen	20.61
Nitrogen	77.95
Carbonic acid gas	.04
Aqueous vapour	1.40
Nitric acid	} traces.
Ammonia	
Carburetted hydrogen	} traces in towns.
Sulphuretted hydrogen	
Sulphurous acid	

The above proportions are by volume. By weight there will be nearly 23 parts of oxygen to nearly 77 of nitrogen, oxygen being the heavier in the ratio of 16 to 14. Ozone (*vide* Index) is also said to be a normal constituent of air.

The comparative inactivity or negative character of nitrogen in its free condition,—that is when uncombined with other elements,—contrasts strongly with its apparent influence in a state of combination. When its compounds with hydrogen come to be studied, it will be found to be, apparently, the chief, or leading, or, in a sense, the most important element of those compounds—the ammoniacal compounds. United with carbon, it gives the poisonous cyanic substances. With oxygen, it yields quite a large group of bodies, amongst which are the common and important class of salts termed nitrates. With carbon as well as hydrogen and some oxygen, it affords powerful agents termed alkaloids—near relatives of ammonia; while the same elements otherwise grouped, and sometimes a little sulphur or phosphorus, form the various albumenoid and gelatigenous matters characteristic of the tissues of animals and vegetables. In a perfect structure we should perhaps scarcely regard any one element or member as more important than another, still such a conclusion almost forces itself upon us as we become acquainted with the chemical history of combined nitrogen.

CHLORINE.

Source.—This element is a gas. Its chief source is common salt, more than half of which is chlorine.

Preparation.—About a quarter of an ounce of salt and the same amount of black oxide of manganese are mixed, and placed in a test-tube with sufficient water to cover them; on adding a small quantity of sulphuric acid, the evolution of chlorine commences. For mode of collection see following paragraphs.

Another process.—As the action of the sulphuric acid on the salt in the above process is mainly to give hydrochloric acid, the latter acid (about 4 parts) and the black oxide of manganese (about 1 part) may be used in making the gas, instead of salt, sulphuric acid, and black oxide of manganese.

This, the usual process, is that adopted in the "British Pharmacopœia."

Fig. 10.

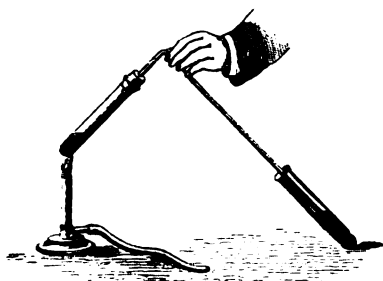
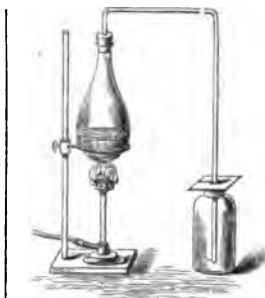


Fig. 11.



PREPARATION OF CHLORINE.

Collection and Properties.—Chlorine is a most suffocating gas. Great care must consequently be observed in experimenting with this element. As soon as its penetrating odour indicates that it is escaping from the test-tube, the cork and delivery-tube (similar to that used in making oxygen) should be fitted on, and the gas allowed to pass to the bottom of another test-tube containing water (fig. 10). When thirty or forty small bubbles have passed, their evolution being assisted by slightly heating the generating-tube, the latter should be removed to the cupboard usually provided in laboratories for performing operations with noxious gases, or dismantled, and the contents washed away. The water in the collecting-tube will now be found to smell of the gas, chlorine being, in fact, soluble in about half its bulk of water. Chlorine-water is official* in the "British Pharmacopœia" (*Liquor Chlori*).

* The Pharmacopœia and all in it is official (*officæ*, Fr. from *L. officium*, an office). There are many things which in pharmacy are officinal (Fr. from *L. officina*, a shop) but not official. To restrict the word *official* to the contents of a pharmacist's shop, and to that portion of the contents which is Pharmacopœial, is radically wrong, and should be avoided.

"An *official* formula is one given under authority. An *officinal* formula

Larger quantities may be made from hydrochloric acid and black oxide of manganese (4 to 1) in a florence flask fitted with a delivery-tube, the flask being supported over a flame by the ring of a retort-stand or any similar mechanical contrivance (fig. 11). A piece of cardboard on the neck of the collecting-bottle, as indicated in the figure, retards diffusion of the gas from the bottle during collection of the gas.

Mem.—Flasks and similar glass vessels are less liable to fracture if protected from the direct action of the flame by being placed on a piece of wire gauze 2 to 4 inches square, or on a *sand-bath*, that is, a saucer-shaped tray of sheet iron on which a thin layer of sand is placed.

The *Vapor Chlori*, B. P., or Inhalation of Chlorine, is simply moist chlorinated lime so placed that some of the chlorine given off may be inhaled.

During these manipulations the operator will have noticed that chlorine is of a light green colour. The tint is readily observed when the gas is collected in large vessels. As it is soluble in water ($2\frac{1}{2}$ vols. in 1 vol. at 60° F.), it cannot be economically stored over that liquid. Being, however, nearly twice and a half as heavy as air, it may be collected by simply allowing the delivery-tube to pass to the bottom of the test-tube or dry bottle (fig. 11).

The distinctive property of chlorine is its bleaching-power. Prepare some coloured liquid by placing a few chips of logwood or other dyeing-material in a test-tube half full of hot water. Pour off some of this red decoction into another tube, add a few drops of the chlorine-water; the red colour is rapidly destroyed.

Chlorine readily decomposes offensive effluvia; it is one of the most powerful of the *deodorizers*. It also decomposes putrid and infectious matter; it is one of the best of *disinfectants*.

is one made in obedience to the customary usage of the shop (*officina*). To state that any preparation under the sanction of the British Pharmacopœia is officinal, is a misapprehension of the meaning of the word."—J. BAUGH.

"That is *official* which emanates from a recognised authority. That is *officinal* which is issued from an *officina* or workshop."—J. INCE.

"*Official* writings and orders are those issued by official persons. *Officinal* articles are such as are found in a shop."—J. F. STANFORD, M.A., F.R.S.

(*Antiseptics* are substances which *prevent* putrefaction. See Index.)

Combination of Hydrogen with Chlorine, forming Hydrochloric Acid.—If an opportunity occurs of generating chlorine in a closed chamber or in the open air, a test-tube, of the same size as one of those in which hydrogen has been retained from a previous operation, is filled with the gas. The hydrogen-tube is then inverted over that containing the chlorine, the mouths being kept together by encircling them with a finger. After the gases have mixed, the mouths of the tubes are quickly in succession brought near a flame, when explosion occurs, and fumes of a compound of hydrochloric acid with the moisture of the air are formed. The Hydrochloric Acid of pharmacy is a solution of the gas (made in a more economical way) in water.

The foregoing experiment affords evidence of the powerful affinity of chlorine and hydrogen for each other. Chlorine dissolved in water will, in sunlight, slowly remove hydrogen from some of the water and liberate oxygen. The bleaching-power of chlorine is generally referred to this oxidizing effect which it produces in presence of water; for dry chlorine does not bleach.

Density.—Chlorine is thirty-five and a half times as heavy as hydrogen. A wine-bottle would hold about $35\frac{1}{2}$ grains.

SULPHUR, CARBON, IODINE.

The physical properties of these elements (colour, hardness, weight, etc.) are familiar. Their leading chemical characters will also be understood when a few facts concerning each are made the subject of experiment.

SULPHUR.—Burn a small piece of sulphur; a penetrating odour is produced, due to the formation of a colourless gas, the same as that formed on igniting a sulphur-tipped lucifer match.

This product is a perfectly definite chemical compound of the oxygen of the air with the sulphur. It is termed sulphurous anhydride or sulphurous acid gas.

CARBON is familiar in the forms of soot, coke, charcoal, graphite (or plumbago, popularly termed blacklead), and diamond. The presence of carbon in wood, and in other vegetable and animal matter, is at once rendered evident by heat. Place a little tartaric acid on the end of a knife in a flame; the blackening that occurs is due to the separation of carbon. The black matter at the extremity of a piece of half-burned wood is also carbon.

Carbon, like hydrogen, phosphorus, and sulphur, has a great affinity for oxygen at high temperatures. A striking evidence of that affinity is the evolution of sufficient heat to make the materials concerned red- or even white-hot. When ignited in the dilute oxygen of the air, carbon simply burns with a moderate glow, as seen in an ordinary coke or charcoal fire; but when ignited in pure oxygen, the intensity of its combination is greatly exalted. The product of the combination of the two elements, if the oxygen be in excess, is an invisible gaseous body termed carbonic acid gas; if the carbon be in excess, another invisible gas termed carbonic oxide results.

IODINE.—A prominent chemical characteristic of iodine is its great affinity for metals. Place a piece of iodine, about the size of a pea, in a test-tube with a small quantity of water, and add a few iron filings or small nails. On gently warming this *mechanical* mixture, or even shaking if longer time be allowed, the colour and odour of the iodine disappear; it has *chemically* combined with the iron, a *chemical compound* has been produced. If the solution be filtered, a clear aqueous solution of the compound of the two elements is obtained.

This compound is an iodide of iron. Its solution, made as above, and mixed with sugar, forms, when of a certain strength, the ordinary Syrup of Iodide of Iron of pharmacy (*Syrupus Ferri Iodidi*, B. P.) A strong solution mixed with sugar and liquorice-root constitutes the corresponding Pill (*Pilula Ferri Iodidi*, B. P.). The solid iodide (*Ferri Iodidum*, B. P.) is obtained on removing the water of the above solution by evaporation.

Sulphur and Iron, also, when very strongly heated, *chemically combine* to form a substance which has none of the properties

of a *mixture* of sulphur and iron—that is, has none of the characters of sulphur and none of iron, but new properties altogether. The product is termed Sulphide of Iron. Its manufacture and uses will be alluded to in treating of the compounds of iron; it is mentioned here as a simple but striking illustration of the difference between a *chemical compound* and a *mechanical mixture*.

THE ELEMENTS, THEIR SYMBOLS, ETC.

From the foregoing statements a general idea will have been obtained of the nature of several of the more frequently occurring elements. Some additional facts concerning them may be gathered from the following Table, which gives the name in full, the symbol (or short-hand character*) of the name, and its origin.

For the purposes of study the elements may be divided into three classes, viz., those frequently used in pharmacy, those seldom, and those never used.

NAME.	SYMBOL.	DERIVATION OF NAME.
Oxygen . . .	O	From ὀξύς (oxūs) <i>acid</i> , and γένεσις (genesis) <i>generation</i> , i.e. generator of acids. It was supposed to enter into the composition of all acids when first discovered.
Hydrogen . .	H	From ὑδὼρ (hudōr) <i>water</i> , and γένεσις (genesis) <i>generation</i> , in allusion to the product of its combustion in air.
Nitrogen . .	N	From νίτρον (nitron), and γένεσις (genesis), <i>generator of nitre</i> .
Carbon . . .	C	From carbo, <i>coal</i> , which is chiefly carbon.
Chlorine . .	Cl	From χλωρός (chlōros) <i>green</i> , the colour of this element.
Iodine . . .	I	From ἰώ (ion) <i>a violet</i> , and εἶδος (eidos) <i>likeness</i> , in reference to the colour of its vapour.
Sulphur . . .	S	From sal <i>a salt</i> , and πῦρ (pūr) <i>fire</i> , indicating its combustible qualities. Its common name, <i>brimstone</i> , has the same meaning, being the slightly altered Saxon word <i>brynstone</i> , i.e. <i>burnstone</i> .

* The symbol is also much more than the short-hand character, as will presently be apparent.

NAME.	SYMBOL.	DERIVATION OF NAME.
Phosphorus .	P	Φῶς (phōs) <i>light</i> , and φέρειν (pherein) <i>to bear</i> . The light it emits may be seen on exposing it in a dark room.
Potassium . . (Kalium.)	K	<i>Kalium</i> , from <i>kali</i> , Arabic for <i>ashes</i> . Manufactories in which certain compounds of potassium and allied sodium-salts are made are called <i>alkali-works</i> to this day. <i>Potassium</i> , from <i>pot-ash</i> ; so called because obtained by evaporating the lixivium of wood-ashes in pots. From such ashes the element was first obtained, hence the name.
Sodium . . . (Natrium.)	Na	<i>Natrium</i> , from <i>natron</i> , the old name for certain natural deposits of carbonate of sodium. <i>Sodium</i> , from <i>soda-ash</i> or <i>sod-ash</i> , the residue of the combustion of masses or <i>sods</i> of marine plants. These were the sources of the metal.
Ammonium .	Am (NH ₄)	This body is not an element; but its components exist in all ammoniacal salts, and apparently play the part of such elements as potassium and sodium. Sal ammoniac (chloride of ammonium) was first obtained from near the temple of Jupiter Ammon in Libya; hence the name.
Barium . . .	Ba	From βαρύς (barūs) <i>heavy</i> , in allusion to the high specific gravity of "heavy spar," the most common of the barium minerals.
Calcium . .	Ca	<i>Calx</i> , <i>lime</i> , the oxide of calcium.
Magnesium .	Mg	From <i>Magnesia</i> , the name of the town (in Asia Minor) near which the substance now called "native carbonate of magnesia" was first discovered.
Iron (Ferrum.)	Fe	The spelling is from the Saxon <i>iren</i> , the pronunciation probably from the kindred Gothic " <i>iarn</i> ." The derivation is Aryan; it probably originally meant <i>metal</i> .
Aluminium .	Al	The metallic basis of alum was at first confounded with that of sulphate of iron, which was the alum of the Romans, and was so called in allusion to its tonic properties, from <i>alo</i> , <i>to nourish</i> .
Zinc	Zn	From Ger. <i>Zinn</i> , <i>tin</i> , with which zinc seems at first to have been confounded.
Arsenicum .	As	Ἀρσενικόν (arsenikon), the Greek name for orpiment, a sulphide of arsenicum. Common white arsenic is an oxide of arsenicum.

NAME.	SYMBOL.	DERIVATION OF NAME.
Antimony . . . (Stibium.)	Sb	$\Sigma\tau\beta\iota$ (stibi), or $\sigma\tau\iota\mu\mu$ (stimmi) was the Greek name for the native sulphide of antimony. The word <i>antimony</i> is said to be derived from $\acute{\alpha}\nu\tau\iota$ (anti) <i>against</i> , and <i>moine</i> , French for <i>monk</i> , from the fact that certain monks were poisoned by it. From <i>Cyprus</i> , the name of the Mediterranean island where this metal was first worked.
Copper . . . (Cuprum.)	Cu	The Latin word is expressive of "something heavy," and the Saxon <i>led</i> has a similar signification.
Lead . . . (Plumbum.)	Pb	<i>Hydrargyrum</i> , from $\eta\delta\omega\rho$ (hudōr) <i>water</i> , and $\acute{\alpha}\rho\gamma\upsilon\rho\omicron\varsigma$ (arguros) <i>silver</i> , in allusion to its liquid and lustrous characters. <i>Mercury</i> , after the messenger of the gods, on account of its susceptibility of motion. The old name <i>quicksilver</i> also indicates its ready mobility and argentine appearance.
Mercury . . . (Hydrargyrum.)	Hg	<i>Argyros</i> (arguros) <i>silver</i> , from $\acute{\alpha}\rho\gamma\upsilon\varsigma$ (argos) <i>white</i> . Words resembling the term <i>silver</i> occur in several languages, and indicate a white appearance.
Silver . . . (Argentum.)	Ag	

The following are names of some of the less frequently occurring elements, compounds of which, however, are alluded to in the British Pharmacopœia, or are met with in pharmacy.

Bromine . . .	Br	From $\beta\rho\acute{\omicron}\mu\omicron\varsigma$ (brōmos), a <i>stink</i> . It has an intolerable odour.
Fluorine . . .	Fl	From <i>fluo</i> , to <i>flow</i> . Fluoride of calcium, its source, is commonly used as a flux in metallurgic operations.
Boron . . .	Bo	From <i>borak</i> or <i>baurak</i> , the Arabic name of <i>borax</i> , the substance from which the element was first obtained.
Silicon . . .	Si	From <i>silex</i> , Latin for <i>flint</i> , which is nearly all silica (an oxide of silicon).
Lithium . . .	L	From $\lambda\iota\theta\epsilon\iota\omicron\varsigma$ (litheios) <i>stony</i> , in allusion to its supposed existence in the mineral kingdom only.
Strontium . . .	Sr	This name is commemorative of <i>Strontian</i> , a mining-village in Argyleshire, Scotland, in the neighbourhood of which the mineral known as <i>strontianite</i> or carbonate of strontium was first found.

NAME.	SYMBOL.	DERIVATION OF NAME.
Cerium . . .	Ce	Discovered in 1803, and named after the planet <i>Ceres</i> , which was discovered on Jan. 1, 1801. The oxalate of cerium is official, but seldom used.
Chromium . .	Cr	From <i>χρῶμα</i> (<i>chrōma</i>) <i>colour</i> , in allusion to the characteristic appearance of its salts.
Manganese . .	Mn	Probably a mere transposition and repetition of most of the letters of the word <i>magnesia</i> , with whose compounds those of manganese were confounded till the year 1740.
Cobalt . . .	Co	<i>Cobaltus</i> or <i>Kobold</i> was the name of a demon supposed to inhabit the mines of Germany. The ores of cobalt were formerly troublesome to the German miners, and hence received the name their metallic radical now bears.
Nickel . . .	Ni	<i>Nickel</i> , from <i>nil</i> , is a popular German term for <i>worthless</i> . The mineral now known as nickel ore was formerly called by the Germans <i>Kupfernickel</i> , <i>false copper</i> , on account of its resemblance to copper (<i>Kupfer</i>) ore. When a new metallic element was found in the ore, the name nickel was retained.
Tin (Stannum).	Sn	Both words are possibly corruptions of the old British word <i>staen</i> , or the Saxon word <i>stan</i> , a stone. Tin was first discovered in Cornwall, and the ore (an oxide) is called <i>tinstone</i> to the present day.
Gold (Aurum) .	Au	<i>Aurum</i> (Latin) from a Hebrew word signifying the colour of fire.
Platinum . .	Pt	<i>Gold</i> ; a similar word is expressive of <i>bright yellow</i> in several old languages.
Bismuth . .	Bi	From <i>platina</i> (Spanish), diminutive of <i>plata</i> , <i>silver</i> . It somewhat resembles silver in appearance, but is less white and lustrous.
Cadmium . .	Cd	Slightly altered from the German <i>Wismuth</i> , derived from <i>Wiesematte</i> "a beautiful meadow," a name given to it originally by the old miners in allusion to the prettily variegated tints presented by the freshly exposed surface of this crystalline metal.
		<i>Kadmeia</i> (Kadmeia) was the ancient name of calamine (carbonate of zinc), with which carbonate of cadmium was long confounded, the two often occurring together.

Gold, Platinum, Tin, and Silicon are classed with the less important elements, because their salts are seldom used in pharmacy.

It will be noticed that the symbol of an element is simply the first letter of its Latin name, which is generally the same as in English. Where two names begin with the same letter, the less important has an additional letter added.

QUESTIONS AND EXERCISES.

1. Of how many elements is terrestrial matter composed ?
2. In what state do the elements occur in nature ?
3. Distinguish between the *art* and the *science* of chemistry.
4. What is the difference between an element and a compound ?
5. Enumerate the chief non-metallic elements.
6. Describe a process for the preparation of oxygen.
7. How are gases usually stored ?
8. Mention the chief properties of oxygen.
9. What is the source of animal warmth ?
10. State the proportion of oxygen in air.
11. Is the proportion constant, and why ?
12. Give a method for the elimination of hydrogen from water.
13. State the properties of hydrogen.
14. Why is a mixture of hydrogen and air explosive ?
15. Explain the effects producible by the ignition of large quantities of coal-gas and air.
16. What is the nature of combustion ?
17. Define a *combustible* and a *supporter of combustion*.
18. Describe the structure of flame.
19. State the principle of the Davy safety-lamp.
20. To what extent is hydrogen lighter than oxygen ?
21. What do you mean by *diffusion* of gases ?
22. State Graham's law concerning diffusion.
23. Name the source of phosphorus, and give its characters.
24. Why does phosphorus burn in air ?
25. What remains when ignited phosphorus has removed all the oxygen from a confined portion of air ?
26. Mention the properties of nitrogen.
27. What office is fulfilled by the nitrogen of air ?

28. State the proportions of the chief constituents of air.
 29. Mention the minor or occasional constituents of air.
 30. What is the proportion by weight of nitrogen to oxygen in the atmosphere?
 31. Give the specific gravity of nitrogen.
 32. How is chlorine prepared?
 33. Enumerate the properties of chlorine.
 34. Define the terms *deodorizer* and *disinfectant*.
 35. Explain the bleaching effect of chlorine.
 36. What proportion of hydrogen to chlorine is necessary for the formation of hydrochloric acid gas?
 37. State the prominent chemical and physical characters of sulphur.
 38. State the prominent characters of carbon.
 39. State the prominent characters of iodine.
 40. Give the derivations of the names of some of the elements.
 41. What are the symbols of oxygen, hydrogen, nitrogen, carbon, chlorine, iodine, sulphur, phosphorus?
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THE LEARNER IS RECOMMENDED TO READ THE FOLLOWING PARAGRAPHS ON THE GENERAL PRINCIPLES OF CHEMICAL PHILOSOPHY CAREFULLY ONCE OR TWICE, THEN TO STUDY (EXPERIMENTALLY, IF POSSIBLE) THE SUCCEEDING PAGES, RETURNING TO AND READING OVER THE GENERAL PRINCIPLES FROM TIME TO TIME UNTIL THEY ARE THOROUGHLY COMPREHENDED.

Note.—Throughout this Manual paragraphs describing experiments to be performed are distinguished from paragraphs containing matter merely to be read by being printed in somewhat more widely spaced type.

THE GENERAL PRINCIPLES OF CHEMICAL PHILOSOPHY.

DEFINITION OF CHEMICAL ACTION.

The learner may now proceed to study the manner in which substances act chemically on each other. By acting *chemically* it will be obvious, from the preceding experiments, that what is meant is *so affecting each other that the substances are greatly altered in properties*. A mixture of oxygen and hydrogen is still a gas; a chemical compound of oxygen and hydrogen is a liquid, namely, water: here is a great alteration in leading properties. Iodine is only slightly soluble in water, and forms a brown coloured solution, and iron is insoluble; but when iodine and iron are *chemically* combined, the product is very soluble in water, forming a light green solution in which the eye can detect neither iodine nor iron, and which is utterly unlike iron or iodine in any one of their properties. Sand, sugar, and butter rubbed together form a mere mixture, from which water would extract the sugar, and ether dissolve out the butter, leaving the sand. Tartaric acid, carbonate of sodium, and water added to each other form a *chemical compound* containing neither tartaric acid nor carbonate of sodium, these bodies having attacked each other and formed fresh combinations. These illustrations show that *chemical action* is distinguished by (*a*) producing an entire change of properties in bodies. It also is (*b*) exerted only between definite weights and volumes of matter. This (*a* and *b*) cannot be said of any other action—the action of any of the other great forces of nature (gravitation, heat, light, electricity, etc.); hence the statements (*a* and *b*) furnish a sharp and precise definition of *chemical action* or the chemical force, the force whose manifestations the reader of this book, or of any other manual of chemistry, is studying.

ATOMS.

In a chemical compound, what has become of its constituents? Let the reader place before him specimens of sulphur, iron, and sulphide of iron; or iodine, iron, solid iodide of iron, and its solution in water or syrup (*Syrupus Ferri*

Iodidi, B. P.). In the sulphide of iron, what has become of the sulphur and of the iron from which it was made? The mixture of sulphur and iron in combining to form sulphide of iron has not lost weight, and, indeed, by certain processes it is possible to recover its sulphur as sulphur, and its iron as iron; so that we are compelled to believe (we cannot avoid the conclusion) that sulphide of iron contains particles of sulphur and of iron. But how small must be these particles! Rub a minute fragment to dust in a mortar and place a trace of the powder under the highest power of the best microscope; no yellow particle is visible, not the minutest portions of lustrous metal, but dull brown miniature fragments of the original mass. The elementary particles of sulphur and iron, or of the elements in any other compound (the chlorine and sodium in common salt, or the iodine and iron in solution of iodide of iron), are, in short, too small to be seen. Can they be imagined? Again, no! The mind cannot conceive of a particle of any thing (sulphur, iron, sulphide of iron, or what not) so small but what the next instant the imagination has divided it. Yet learner and teacher must have some common platform on which to reason and converse. The difficulty is met by speaking of these inconceivably small particles as *atoms* (*ἄτομος*, *atomos*, indivisible; from the privative *a* and *τέμνω*, *temnō*, to cut—that which is not cut, or divided), an expedient suggested by our countryman Dalton at the commencement of the present century. It is an expedient not, perhaps, altogether satisfactory, but is the only one possible to the majority of minds in the present state of knowledge and education. We cannot speak of iodine and iron uniting lump to lump, as two bricks are cemented together or blocks of wood glued together, for such is not the kind of action. We cannot select minute fragments of each to regard as the combining portions; for the minutest fragment we could obtain is visible, and iodide of iron contains neither visible iodide nor visible iron. And yet iodide of iron contains both iodine and iron, or, at least, a given weight of the compound is obtained from the same weight of constituents, and the same weight of constituents is obtainable from an equal weight of the compound. We might say that molecules are concerned in the operation; but molecules means little masses of—of what! there is positively no word left with which to carry on conversation and description but *atoms*. Any other mode of treating the matter is too subjective for general employment.

Moreover, any difficulty in forming a definite conception of an atom is met by regarding an atom, not necessarily as something which *cannot* be divided, but as "a particle of matter which undergoes no further division in *chemical metamorphoses*" (Kekulé). Even physicists regard atoms from much the same point of view; indeed, they often speak of still larger portions of matter (molecules) as atoms, meaning thereby "something which is not divided in *certain cases that we are considering*" (Clifford).

THE CHEMICAL FORCE.

What power binds the atoms of a chemical compound together in such marvellous closeness of union that in the couple or group they lose all individuality? Clearly an attractive force of enormous power, a force remotely resembling, perhaps, that which attracts a piece of iron to a magnet. Only by such an assumption can we conceive that common salt contains chlorine and a metal (sodium), or that wood contains carbon, hydrogen, and oxygen. Were not this force thus all-powerful, the carbon in wood would show its blackness and other qualities, and the hydrogen and oxygen give indications of their gaseous and other characters. This attractive force is commonly termed *the chemical force*, sometimes *chemical affinity*. The word *chemism* has also been proposed for it, just as the magnetic force is termed magnetism, but has not been generally adopted.

MOLECULES.

A free, uncombined atom probably cannot exist in a state of isolation at common temperatures for any appreciable length of time. For we must regard an atom as the home of an attractive force of great intensity, and the moment such an atom is liberated from a state of combination (say, hydrogen from water, or chlorine from salt) it finds itself in proximity to another atom having similar desires for union, so to speak; the result is an impetuous rushing together and formation of either couples, trios, or groups, according to the nature of the atoms. It would be as difficult to conceive of separate atoms as to imagine that a strong magnet and a piece of steel could be suspended close to each other without being drawn together. It is, doubtless, possible to keep some pairs of atoms apart by the aid of heat, just as the magnet and steel may be parted by

a superior amount of force, but such a condition of things is probably abnormal. These pairs and other groups of atoms are conveniently designated by the one word *molecule*, the diminutive of *mole*, a mass; literally, little masses. Dissimilar kinds of atoms seem to have greater attraction for each other than similar kinds; for, first, the masses of matter met with in nature in the great majority of cases contain two or more dissimilar elements; and, secondly, at the moment certain elements are liberated from their combination, they are very specially active in combining with other, different, elements; that is to say, the chances are not equal that the liberated elements will either retain their elementary condition or combine to form compounds, but the cases in which compounds are formed are actually in great majority.

RECAPITULATION.

It is desirable that the learner should here make some experiment which will serve to bring again under notice in an applied or concrete form what has just been stated respecting the substances termed *chemical compounds*, and concerning the character of that *chemical force* which resides in the *atoms of molecules*. The following will usefully serve this purpose; it is the process for detecting a trace of sulphurous acid in common liquid hydrochloric acid.

As already proved, hydrogen gas and chlorine gas, when united, form hydrochloric acid gas: the latter dissolved in water is the ordinary liquid of the shops termed *Hydrochloric Acid*, the *Acidum Hydrochloricum* of Pharmacopœias. Commercial samples of this liquid not unfrequently contain as an impurity a trace of sulphurous acid gas, a body also already mentioned and experimentally prepared—a trace too small to be detected by its odour. Obtain a specimen of common liquid hydrochloric acid containing as an impurity a trace of sulphurous acid, or adopt the more simple course of purposely adding a few drops of aqueous solution of sulphurous acid (*Acidum Sulphurosum*,* B. P.) to some hydrochloric acid. (If no sulphurous acid is at hand, the object may be accom-

* These aqueous solutions of acids are generally, for the sake of brevity, simply termed acids.

plished by pouring a quarter or half an ounce of liquid hydrochloric acid into a wide-mouthed bottle, then burning a fragment of sulphur on a wire or strip of wood inside the bottle for a few seconds, and shaking the gas and liquid together.) Pour some of the impure liquid hydrochloric acid into a test-tube, add about an equal bulk of water, and then drop in some fragments of the metal zinc. Effervescence will occur, due to the escape of inodorous hydrogen gas, together with a small quantity of a badly-smelling gas termed sulphuretted hydrogen. Bring the mouth of the tube under the nose; the presence of sulphuretted hydrogen will at once be recognised.

The hydrochloric acid has now been *tested* for sulphurous acid. If the experiment be performed on any commercial specimen of the acid, and a smell of sulphuretted hydrogen be observed, the operator will at once be able to state that the specimen contains sulphurous acid as an impurity.

Using Dalton's theory of the atomic constitution of matter, the explanation of what occurs in the successive steps of the foregoing experiment, is as follows:—

Hydrochloric acid is a *chemical compound* of hydrogen and chlorine. That it is a chemical compound, and not a mere mechanical mixture of hydrogen and chlorine, is shown by the fact that its properties are altogether different from the properties of its constituents. The attractive power or *chemical force* resident in the *atoms* of chlorine and of hydrogen has caused them to combine in the closest manner imaginable and form pairs of atoms or *molecules* of the chemical compound—hydrochloric acid. Zinc being introduced into the acid, and the atoms of zinc and chlorine having even still greater attraction for each other than the hydrogen for the chlorine, the zinc and chlorine atoms combine and form a new molecule (termed chloride of zinc) which remains in the liquid, while the hydrogen atoms, having the atoms of no other element to combine with if the acid is pure, unite to form pairs, or molecules, of hydrogen, and in that state escape from the vessel. If the acid be impure from the presence of sulphurous acid (sulphurous acid gas, it will be remembered, is a compound of sulphur and oxygen), some of the hydrogen atoms, at the moment of birth, their *nascent* state (from *nascor*, to be born)—the specially active state—finding the atoms of

other elements present, namely, the atoms of sulphur and oxygen of the sulphurous acid molecules, combine by preference with these atoms and form new molecules, the sulphur and hydrogen forming sulphuretted hydrogen, and the oxygen and hydrogen producing water—the former escapes with the great bulk of the hydrogen, while the water remains with the water already in the vessel.

Note.—Ordinary hydrogen gas—that is, hydrogen not in the nascent condition—will not thus attack sulphurous acid. Doubtless the amount, or extent of attraction of two atoms of hydrogen for one atom of, say, the sulphur in the sulphurous acid molecule, is a constant amount; but the uncombined, nascent atoms can, it is only fair to suppose, get much nearer to the attacked molecule than they can after they have themselves combined to form a molecule, molecules (but not atoms) having an appreciable amount of space between them, as will be further shown almost immediately. In other words, it is probably distance which prevents an attack which would be inevitable at close quarters. These remarks apply to all similar reactions of other elements.

CONDITIONS AND NATURE OF THE MANIFESTATION OF THE CHEMICAL FORCE.

The exertion of chemical affinity is only possible when the masses of the bodies touch. Thus it was necessary to bring the oxygen, hydrogen, phosphorus, chlorine, sulphur, carbon, iodine, and iron into ordinary contact, in the respective experiments with those elements, before the various reactions occurred. The exact nature of these actions, as indeed of all in which substances act chemically, would seem to be an interchange, most generally a mutual one, of the atoms of which the molecules consist—a change of partners, so to speak. Thus in the experiment in which hydrogen and chlorine gases united to form hydrochloric acid gas, a pair of atoms in a hydrogen molecule and a pair of atoms in a chlorine molecule, finding themselves opposite to each other, changed places, the atoms of each of the old molecules unlinking, so to say, and pairing off in fresh couples—as two brothers who for many years have been close companions, and two sisters similarly united, thrown freshly into each other's society, soon accept new and still more congenial companionship.

{ Hydrogen } and { Chlorine } be- { Hydrogen } and { Hydrogen }
{ Hydrogen } and { Chlorine } come { Chlorine } and { Chlorine }

Or, using the symbols of these elements instead of the full names,

H H and Cl Cl become H Cl and H Cl.

Still further economizing space and trouble, the same statement may be made in the following form :—

H_2 and Cl_2 become $2HCl$.

Once more, by using the *plus* sign (+), instead of the words “and” or “added to,” and the sign or symbol = or *equal*, instead of the words “become” or “are equal to,” we reach the shortest method of expressing this chemical action :—

$H_2 + Cl_2 = 2HCl$

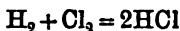
This is the form in which such an action may be expressed in the student's note-book. It is the shortest and most convenient form, and is instructive and suggestive to the mind.

CHEMICAL NOTATION.

We have thus gradually arrived at a spot in the path of chemical philosophy at which we must halt to more fully discuss the usual method of recording chemical travels. We have arrived at the subject of *chemical notation* (from *noto*, I mark), the art or practice of recording chemical facts by short marks, letters, numbers, or other signs. Already the first capital letter, or the first and one of the following small letters of the Latin names of the elements have been employed as contractions, or short-hand expressions, or *symbols* of the whole name. Thus H has been used for the word “hydrogen,” and Cl for “chlorine.” A *second function* of such a symbol is that of indicating one *atom*. Thus H stands not only for the word or substance “hydrogen,” but for one atom of hydrogen. Large and small figures (2 or ₂) indicate a corresponding number of atoms, the small figure only multiplying the one particular symbol to which it is attached, while a larger figure multiplies all the symbols it precedes. Thus H_2 means two atoms of hydrogen, and Cl_2 two atoms of chlorine; while $2HCl$ means two atoms of hydrogen and two atoms of chlorine, or, in one word, two molecules of hydrochloric acid gas. A *third function* of such a symbol as H or Cl is that of indicating one *volume* of the element in the gaseous state. Thus H, Cl, or O, stand, first, for the sub-

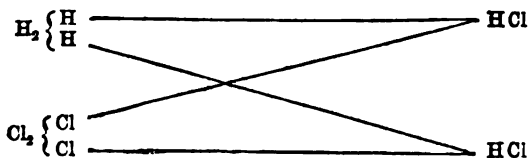
stances named hydrogen, chlorine, and oxygen; secondly, for single atoms of hydrogen, chlorine, and oxygen; thirdly, they represent single and equal volumes of chlorine, hydrogen, and oxygen. It will be remembered that one test-tubeful of hydrogen and an equal sized test-tubeful of chlorine were employed, in a previous experiment, in forming hydrochloric acid gas, HCl.

The *position* of symbols counts for something. Thus HCl indicates not only the substances hydrogen and chlorine, single atoms of each of the substances, and equal volumes of each, but also that the two substances are *joined together by the chemical force*. If the two letters were placed one under the other, or at some distance apart, or were separated by a comma or a plus sign (+), they would be understood to mean a mere mixture of the elements; but placed as close as the printer's types will conveniently and consistently allow, they must be considered to stand for a compound of the elements, that is to say, hydrochloric acid gas (HCl). The collection of symbols representing a molecule is termed a *formula*. H_2 , Cl_2 , and HCl are the *formulae* of hydrogen, chlorine, and hydrochloric acid gas.



Such a set of letters, figures, and marks as that on the above line is collectively termed an *equation*, because it indicates the equality of the number and nature of the atoms before and after chemical action. On the left-hand of the sign of equality (=) are shown two molecules, and on the right-hand two molecules; but, of the molecules on the left, one contains two atoms of hydrogen and the other two atoms of chlorine, while of the molecules on the right each contains one atom of hydrogen and one of chlorine. The equation forms a short and convenient plan of recording the facts of experiment.

Instead of an equation, a *diagram* may be employed to exhibit the same facts. Thus:—



PHYSICAL AND CHEMICAL CONSTITUTION OF
MATTER.

RELATIONS OF GASES, LIQUIDS, AND SOLIDS.

Molecules of gases are not in absolute contact, for a volume of gas may be compressed with very little force to half or one-fourth its bulk—in short, to such an extent that in many cases the molecules sufficiently approximate to form a liquid. In a liquid the molecules are still free to glide about with ease amongst each other; and though in solids they exhibit less mobility, still even solids may be compressed by powerful pressure, so that probably in *no* instance are molecules in absolute contact. (Moreover, from the researches of Caignard de la Tour, and recently of Andrews, there would seem to be no sharp lines of demarcation between the gaseous, liquid, and solid conditions of substances.) One's mental picture of the relative position of the molecules of gaseous, vaporous, liquid, or solid matter must be such a picture as that of the moving particles of dust in the air of a room, or such a relation to each other as that of the planets and stars suspended in space. There is abundant experimental evidence to warrant such a conception. A clear, transparent fluid appears perfectly homogeneous, but is not so. Its particles are not in contact. Every one who has mixed 5 pints of rectified spirit with 3 pints of water knows that the 100 fluid ounces of spirit and 60 fluid ounces of water do not when mixed give 160 ounces of "proof" spirit, but only 156 ounces; the molecules of the liquids have gone closer together, having probably a little attraction for each other. Why a gas under pressure should immediately return to its original bulk when the pressure is removed, while a liquefied or solidified gas only slowly resumes the gaseous or vaporous state, is a question which requires for discussion a knowledge of the nature of forces other than the chemical. For it must be remembered that the study of the chemical force is mainly the study of the internal constitution of molecules, the study of the properties of entire molecules forming the domain of Physics, sometimes termed Natural Philosophy. (Physics, from *φύσις*, *phusis*, nature; that is, visible and material nature; the study of actions and reactions which do not involve entire and permanent change in the properties of bodies—the study of the action of heat, light, electricity, magnetism, gravitation, etc., on matter.)

It is necessary, however, to state something more about the physical as well as the chemical condition of the molecules of a *gas*, in order that the learner may be prepared for the fact that mixtures of certain gaseous elements, in combining to form gaseous compounds, diminish considerably in volume. Thus while a pint of hydrogen and a pint of chlorine give a quart of hydrochloric acid gas,

Hydrogen.	Chlorine.
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Hydrochloric acid gas.

two pints of hydrogen and one of oxygen are necessary to produce a quart of gaseous water (steam). It will be remembered that two volumes of hydrogen and one of oxygen were necessary in a previous experiment in which water was formed.

Hydrogen.	Hydrogen.	Oxygen.
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Gaseous water (Steam).

Now, that a pint of hydrogen gas and a pint of chlorine gas should, after chemical reaction or rearrangement of the atoms of the molecules has taken place, form two pints of hydrochloric acid gas, is quite what we should expect. For, first, the reader, by this time, is not astonished that chemical combination is attended by entire change of properties; and, secondly, the experience of years has led him to expect that a pint of one thing added to a pint of another gives two pints of the mixture. But that two pints of hydrogen and one pint of oxygen should, after combination (and under like conditions of temperature and pressure), give, not three, but two pints of product (steam) is perhaps somewhat astonishing

and needs explanation. To this end let us picture a few of the molecules of hydrogen and as many molecules of chlorine. Draw with a pencil on paper several pairs of crosses (+ +) to represent hydrogen molecules, and circles (°°) for chlorine molecules, or, if coloured ink is at hand, red pairs of dots for hydrogen and green for chlorine. Or, at once, for facility in printing, let the following pairs of letters *hh* represent a few (say, nine) molecules of hydrogen, and *cc* molecules (nine) of chlorine—before combination.

<i>h h</i>	<i>h h</i>	<i>h h</i>	<i>c c</i>	<i>c c</i>	<i>c c</i>
<i>h h</i>	<i>h h</i>	<i>h h</i>	<i>c c</i>	<i>c c</i>	<i>c c</i>
<i>h h</i>	<i>h h</i>	<i>h h</i>	<i>c c</i>	<i>c c</i>	<i>c c</i>

Then, after combination, we shall have eighteen molecules of hydrochloric acid gas:—

<i>h c</i>	<i>h c</i>	<i>h c</i>	<i>h c</i>	<i>h c</i>	<i>h c</i>
<i>h c</i>	<i>h c</i>	<i>h c</i>	<i>h c</i>	<i>h c</i>	<i>h c</i>
<i>h c</i>	<i>h c</i>	<i>h c</i>	<i>h c</i>	<i>h c</i>	<i>h c</i>

But when two volumes of hydrogen and one of oxygen combine and give two volumes of steam, the mental picture must be not that of molecules somewhat nearer to each other than before, nor any difference in the size of the molecules, but a picture of molecules containing three instead of two atoms—thus, still using pairs of letters, just for the moment, to represent a few (the space will allow only twenty-seven) molecules:—

<i>h h</i>	<i>h h</i>	<i>h h</i>	<i>h h</i>	<i>h h</i>	<i>h h</i>	<i>o o</i>	<i>o o</i>	<i>o o</i>
<i>h h</i>	<i>h h</i>	<i>h h</i>	<i>h h</i>	<i>h h</i>	<i>h h</i>	<i>o o</i>	<i>o o</i>	<i>o o</i>
<i>h h</i>	<i>h h</i>	<i>h h</i>	<i>h h</i>	<i>h h</i>	<i>h h</i>	<i>o o</i>	<i>o o</i>	<i>o o</i>

The twenty-seven molecules (eighteen hydrogen, nine oxygen) will, after combination, become eighteen molecules of steam:—

<i>h o h</i>	<i>h o h</i>	<i>h o h</i>	<i>h o h</i>	<i>h o h</i>	<i>h o h</i>
<i>h o h</i>	<i>h o h</i>	<i>h o h</i>	<i>h o h</i>	<i>h o h</i>	<i>h o h</i>
<i>h o h</i>	<i>h o h</i>	<i>h o h</i>	<i>h o h</i>	<i>h o h</i>	<i>h o h</i>

As already suggested, one's mental picture of a number of molecules may well give them such a relation to each other as that of a number of solar systems in the universe, equally distant from each other, and each occupying a similar space, yet one system containing a sun and one planet, another a sun and two planets, and so on, or even one or more of the planets having one or more moons. Indeed, the atoms in some very complex molecules really appear to have very much the relation to each other of the sun, planets, and moons of a solar system. To indicate such molecules by letters as above would of course require more space than is there given to the assumed pictures of molecules. For an amusing and philosophical picture of a complex molecule, the reader is referred to a pamphlet by the late Prof. Clifford entitled "Atoms," by which he means physical atoms, physical units, or what chemists call molecules: it is published by F. Pitman, Paternoster Row, London, and its price is one penny.

Here occurs an opportunity that must not be lost of stating a mode of reasoning by which a molecule of oxygen (or of many other elements) is shown to be a double structure—shown to contain two atoms. Five equal-sized bottles are before us, two filled with hydrogen, one with oxygen, and two with steam. (The bottles are hot enough to prevent the steam condensing to water, and all five are at the same temperature.) Apply heat so that all shall be equally heated, *the three different substances expand equally*. Cool equally, *the contents contract equally*. Apply equal pressure to all five, *each is equally affected*. Diminish pressure equally, *each portion of the three substances equally expands*. Gases (practically steam is a gas, it is simply not a permanent gas)—gases thus similarly affected must be, physically, similarly constructed or constituted (a law which will again be referred to, on page 50); each bottle must contain the same number of particles or molecules, and at any one temperature and pressure the molecules in each must be equally distant from each other. We do not know what actual number or distance, but whatever be the number and distance, it is the same for each bottle. Say that one million is the number, then we shall have a million of molecules in the first hydrogen bottle, a million in the second, a million in the oxygen bottle, and a million in each of the steam bottles. We will cause chemical combination between the two millions of hydrogen molecules and one million of oxygen molecules, producing (as we have seen) two

millions of steam molecules, having the properties already stated. But a molecule of steam contains an atom of oxygen. Hence two millions of steam molecules contain two millions of oxygen atoms, which two millions of oxygen atoms have been obtained from one million of oxygen molecules. Therefore each molecule of oxygen was a double structure—each molecule of oxygen contained two atoms of oxygen. As Clifford says, "you cannot put 50 horses into 100 stables, so that there shall be exactly the same amount of horse in each stable; but you can divide 50 *pairs* of horses among 100 stables."

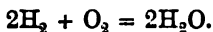
Thus much respecting the constitution of gaseous or vaporous matter. Our knowledge of the constitution of liquid and solid matter is still more limited.

With regard to the notation of the subject, it will be sufficient to state here that while a *symbol* usefully represents one volume of any gas, a *formula of any gas or vapour represents two volumes*. By remembering this general rule we may, by looking at a formula, tell how many volumes of constituents were concerned in the formation of a compound, and therefore what amount of condensation, if any, occurred during the act of formation. By thus reading and interpreting the formula for water, H_2O , we see that two volumes of steam (at any temperature) may be obtained from two volumes of hydrogen and one volume of oxygen (at the same temperature), and thus that the extent of condensation when hydrogen and oxygen (at a stated temperature) unite to form gaseous water (at the same temperature) is from three to two. This subject will again be treated of in connection with those of Chemical Combination and the Specific Gravity of Gases.

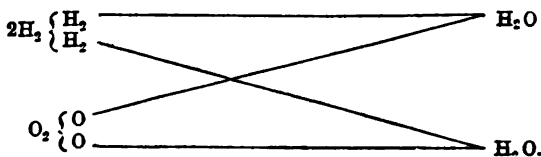
FURTHER REMARKS ON GENERAL CHEMICAL NOTATION.

We may now take an experiment already made as an additional example of chemical action, and describe the simplest way of expressing the same by notation. When two volumes of hydrogen and one of oxygen were caused to combine, the production of flame and noise proved that chemical action of some kind had taken place; had the experiment been performed in dry vessels, evidence of the precise action would have been found in the bedewment or moisture produced by the condensation of the water on the sides of the tube. Similar evidence was afforded on holding a cool glass surface

over the hydrogen-flame. The action is expressed in the following equation:—

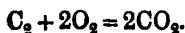


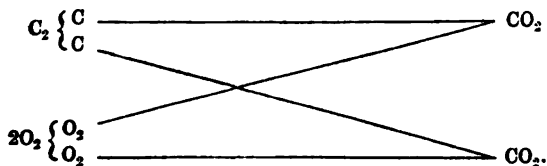
Instead of an equation, the following diagram may be employed:—



The foregoing aggregation of symbols or short-hand characters, or *formula*, H_2O , is, then, a convenient picture of the facts that have already come before us, viz. that water is formed of the elements hydrogen, H, and oxygen, O; moreover, that it is formed of two measures or volumes of hydrogen, H_2 , to one of oxygen, O; and, thirdly, that the molecule of water (H_2O) is formed of two atoms of hydrogen (H_2) and one of oxygen (O). The formula also fulfils the *fourth function* of indicating that the two volumes of hydrogen and one of oxygen in combining condensed to two volumes of steam. That the resulting bulk of steam afterwards shrunk most considerably in condensing to water is another matter altogether, a physical and not a chemical result, and due to the approximation of the molecules of water *after* formation.

Another experiment already performed, illustrating the character of the manifestations of chemical force (symbolically noted as follows), was that in which the red-hot carbon of wood was plunged into oxygen. The evidence of chemical action in that case was the sudden inflammation of the carbonaceous extremity of the wood. The particles of carbon and oxygen having intense attraction or affinity for each other at that temperature, rushed together so impetuously as suddenly to produce a large additional quantity of heat, an amount sufficient to cause the particles to emit an intense white light. The action between carbon and oxygen is expressed on paper in either of the following ways:—





CO_2 is the formula of the well-known gaseous body commonly termed carbonic acid gas.

The reader should here draw for himself equations or diagrams similar to those on pages 36, 42, 43, and thus show the formation of the three other bodies he has already produced—namely, phosphoric anhydride (P_2O_5), sulphurous acid gas (SO_2), and iodide of iron (Fe I_2), submitting the same, if possible, to a tutor or other authority to assure himself of their correctness.

Note.—In the foregoing experiments several illustrations occur of the formation of compounds having the gaseous, liquid, and solid conditions, in one of which three forms all matter in the universe apparently exists.

LAWS OF CHEMICAL COMBINATION (BY WEIGHT).

Chemistry as a science is little more than a hundred years old, though very many of the facts and operations we now term chemical have been known as isolated items of knowledge for centuries. Thus the ancient Egyptians made glass, vitriol, soap, and vinegar; and the Greeks started the idea that matter was composed of a few elements, imagining earth, air, fire, and water to be elements. But the great general principles which interlace and bind together separate facts, those which from their extensive application and importance are denominated *laws*, have all been brought to light since the year 1770.

First Law relating to Chemical Combinations.

Between 1785 and 1800, Bryan Higgins, William Higgins, Wenzel, Richter, and Pronst, made analyses and researches which led up to the following generalization:—*When compounds unite to form definite chemical substances, they always combine in the same proportions.* The curious character of

this fact could but be most striking, and indeed is so now, to the mind receiving it for the first time. Thus water (a compound) added to quicklime (a compound) gives slaked lime, a perfectly definite chemical substance. But whereas sand and water, sugar and water, sand and sugar, and such *mixtures* may be obtained by adding together the ingredients in any proportions whatever, say, 90 of sugar and 10 of sand, or 10 of sugar and 90 of sand, slaked lime (say 100 parts) invariably results from the combination of $75\frac{1}{2}$ of quicklime and $24\frac{1}{2}$ of water. If a larger proportion than $75\frac{1}{2}$ per cent. of quicklime be employed, the excess remains as quicklime mixed with the slaked lime; and if more than $24\frac{1}{2}$ per cent. of water be used, an excess of water remains with the slaked lime and evaporates if the mixture be exposed to the air. Dalton discovered that when *elements* unite to form a definite substance, they, like compounds, always combine in the same proportions; and he was the first to set forth the law in a manner which was at once clear and comprehensive enough to include the former generalization. Thus:—

A definite compound always contains the same elements in the same proportions.

Take another example. Common salt always contains $39\frac{1}{3}$ per cent. of the metal sodium to $60\frac{2}{3}$ of chlorine, and water always 89 of oxygen to 11 per cent. of hydrogen (more exactly 88.89 to 11.11). As with the quicklime and water, so with the chlorine and sodium, and the constituents of many (not *all*) chemical compounds; in such cases, if either be added to the other in any quantity beyond stated proportions, the excess plays no part whatever in the act of combination. (In *some* cases, as will be seen directly, excess of either plays a very simple but very remarkable part.) In short, whether a compound be made directly from its elements, or by the combination of other compounds, or indirectly as one of two products of the action of substances chemically on each other, whatever be its origin, if it is a definite compound it always contains the same elements in the same proportions.

Second Law relating to Chemical Combination.

Dalton further made such experimental researches as enabled him to lay down a second great law. He found that while

many substances only united chemically in one proportion, others combined in two or even more; and he studied several such naturally related bodies. He found that while carbonic oxide (a gas formed when charcoal is burned with an insufficient supply of air) contains such a proportionate weight of carbon and oxygen as is represented by (to use the simplest figures) 3 and 4, carbonic acid (a gas formed when charcoal is burned with excess of air) contains 3 of carbon to exactly twice 4 of oxygen. He proved that a similar relation existed between two compounds of carbon and hydrogen, and between a cluster of compounds of nitrogen and oxygen. The first of the latter, to a given quantity of nitrogen, contains a certain proportion of oxygen; the next, to the same quantity of nitrogen, has exactly twice the proportion of oxygen; and the others have exactly three, four, and five times as much oxygen as the first, the quantity of nitrogen remaining the same throughout. Dalton thus generalized these facts:—

When two elements unite in more than one proportion, the resulting compounds contain, to a constant proportion of one element, simple multiple proportions of the other—or the weights of the constituent elements bear some similar simple relation to each other.

Thus carbonic oxide gas is a definite compound always containing fixed proportions of carbon and oxygen, and carbonic acid gas is also a definite compound always containing fixed proportions of carbon and oxygen. Both thus obey the first law of combination. But whereas carbonic oxide contains, or may be made from, 30 parts (ounces, grains, or other weights) of carbon and 40 of oxygen, carbonic acid contains, or may be made from, 30 parts of carbon and exactly twice 40 of oxygen.

This second law cannot but be as striking as the first when freshly unveiled to the mind. Sand and sugar, or any substances which do not act *chemically* on each other, may be mixed in the proportions of 30 to 40, 30 to 80, 30 to 60, or any other quantities; but if an attempt be made to burn 30 parts of carbon in 60 of oxygen, the elements will themselves naturally assert their own special combining powers, and refuse, so to say, to unite in these proportions: the 30 of carbon will first combine with 40 of oxygen and form 70 of carbonic oxide; and this gas, which, had it the opportunity,

would combine with 40 more of oxygen and form carbonic acid gas, finding only half that quantity, namely 20 of oxygen, present, contents itself by one half (that is 35 of carbonic oxide) accepting the 20 of oxygen and becoming carbonic acid gas, while the other half remains as carbonic oxide. This is a most wonderful fact. Again, if 30 parts of carbon be burnt in more than 80, say 85, of oxygen, only 80 will be used, the other 5 remaining as oxygen merely *mixed* with the resulting carbonic acid gas. If we attempt to burn 30 parts of carbon in less than 40 of oxygen, the oxygen will take up three-fourths its weight of carbon and form carbonic oxide, while the excess of carbon will remain as carbon.

RECAPITULATION.

Nature does not always permit man to mix things in any proportions he pleases. She does sometimes. She does if he only stirs things together, or if he only uses the attractions of adhesion or cohesion in binding the materials together; but if he employs chemical attraction, she restricts him to special proportions. That is to say, if the things mixed do not attack one another or intimately combine, then admixture may be effected in any proportion; and the mixture is a mere mixture having the mean properties of its components. Examples of such mixtures are seen in compound plasters, pill-masses, confections, and plum-puddings. But if the things do unite to form, not a mere mixture having the mean properties of its components, but a compound having new and distinct and definite characters of its own, then nature does not permit man to mix the things in any proportion he pleases. The proportion is one fixed and constant; and if he substitutes proportions of his own, the things unite in the proportions fixed by nature, and the excess he has added either remains in its original uncombined condition, or it combines with the compound already produced to form a second *different* compound. Any one compound, that is, the *same* compound, always contains the same elements in the same proportions, and can only be made from the same elements in the same proportions. An attempt to mix the same elements in other proportions would result in one of two failures, namely, either the extra proportion would remain free and uncombined, or it would combine and convert the first compound, or a por-

tion of it, into a *different* compound. The fresh compound thus produced, like the first, and indeed like all definite compounds, of course always contains the same elements in the same proportions.

In short (law 1), any definite compound always contains the same elements in the same proportions, and (law 2) any two elements uniting in more than one proportion unite in multiples of that proportion and produce so many different definite compounds. Taking hydrogen as uniting in proportions of 1, oxygen unites in proportions of 16—that is 16, twice 16, thrice 16, and so on, never in intermediate proportions. Carbon unites in proportions of 12, sulphur of 32, chlorine $35\frac{1}{2}$. And every other element has its combining proportion fixed by nature.

The student of chemistry is recommended to accept these two great natural facts, great enough to be dignified by the name of *laws*, in all their inherent solidity and simplicity. Of course he will wonder *why* substances should combine, chemically, only in fixed proportions when forming a definite body, and *why*, when a substance combines in more than one proportion to form different definite bodies, the proportions should only be multiple proportions; and an extremely ingenious and useful explanation has been suggested by Dalton (*see* the following paragraphs on the theory that matter is built up of atoms). But man has not yet succeeded in so questioning Nature as to gain from her a *satisfactory* answer to such questions; and until he does succeed, any hypothesis, such as Dalton's, should be held intelligently but loosely. The facts themselves, however, should be grasped with the student's utmost tenacity.

Third Law relating to Chemical Combination.

Careful consideration of the foregoing two great laws has suggested an important truth sometimes termed *The Third Law of Chemical Combination*, namely:—*The proportions in which two elements unite with a third are the proportions (or simple multiples or submultiples of the proportions) in which they unite with each other.* Thus oxygen in proportions of 16 unites with hydrogen, and carbon in proportions of 12 unites with

hydrogen; therefore 16 and 12 are the proportions in which oxygen and carbon will unite with each other.*

THE ATOMIC THEORY.

The laws which Dalton (1803 to 1808) so largely aided to unveil—two grand and wonderful truths—he explained and correlated by a simple and beautiful hypothesis. *Dalton suggested that matter was not infinitely divisible, but composed of minute particles or ATOMS having an invariable character. In the words of Wurtz, "To an old and vague notion he attached an exact meaning by supposing that the atoms of each kind of matter possess a constant weight, and that combination between two kinds of matter takes place not by penetration of their substance, but by juxtaposition of their atoms."*

Thus under this hypothesis, or atomic theory as it is generally termed, carbonic oxide is a definite compound always containing the same elements in the same proportions, *because* each particle of it is composed of an atom of carbon and an atom of oxygen chemically united, the weights of the atoms being in the proportion of 3 and 4; that is, having a constant weight of 12 and 16, as we now believe. Carbonic acid gas is also a definite compound always containing the same elements in the same proportions; and the proportion of oxygen is just double that in carbonic oxide, *because* each particle of it is composed of an atom of carbon (weighing 12), and two atoms of oxygen (each weighing 16).



IMAGINARY PICTURES OF MOLECULES OF CARBONIC OXIDE GAS AND CARBONIC ACID GAS.†

* See Axiom 1 in Hawtrey's fascinating "Introduction to the Elements of Euclid," Longmans & Co., 2s. 6d., a book strongly recommended to any chemical student who is not familiar with the mode of reasoning commonly termed geometrical.

† The size of atoms, their shape, their absolute weight—whether or not they are in actual contact—whether or not they are fixed in relation to each other, free to move about each other, or in a constant state of motion—and whether or not the chemical force actuates them as the force of gravitation influences our earth and moon and solar systems, are matters of which at present we know almost nothing. The two pictures are not intended to convey any impression that the following formulæ do not give:—CO or OC, OCO or OOC, or COO or CO₂.

Again, the facts that with 12 of carbon oxygen unites in the proportion of 16, or multiple of 16; that with 12 of carbon, sulphur unites in the proportion of 32, or a multiple of 32 (the liquid known as disulphide of carbon is a chemical compound of 12 of carbon to twice 32 of sulphur); and thirdly, that oxygen and sulphur unite in proportions of 16 and 32, are at once explained on the assumption that these elements exist in atoms which have the respective weights mentioned. Existing in indivisible particles (atoms) which weigh 16, 12, and 32, oxygen, carbon, and sulphur *must* unite in indivisible weights of 16, 12, and 32.

ATOMIC WEIGHTS.

What has just been stated respecting two or three elements is true of all the elements. It is a fact, that when elements unite with one another in the peculiar and intimate manner termed chemical, they do not combine in the haphazard proportions of a mere mixture, but in one fixed and constant proportion. Such proportions or weights represent, according to Dalton, the weights of their atoms. Oxygen unites with other elements in proportions of 16, therefore 16 is the weight of the atom of oxygen. Chlorine unites with other elements in proportions of $35\frac{1}{2}$, therefore $35\frac{1}{2}$ is the atomic weight of chlorine. And for a similar reason the atomic weights of hydrogen will be 1, carbon 12, sulphur 32, nitrogen 14, and iodine 127. Of course it will be understood that these are the *relative* weights of atoms, for we cannot know the absolute weights. All that is known is that the chlorine atom for instance, is 35.5 times as heavy as the hydrogen atom, whatever the absolute weight of the latter may be, and the iodine atom 127 times as heavy. The quantity of metal which with 35.5 of chlorine will form a chloride, and with twice 35.5 a second chloride (dichloride or bichloride), will require 127 of iodine to form an iodide, and twice 127 of iodine to form a second iodide (a diiodide or biniodide)*.

Note on Notation.—A *fourth function* of a symbol is to represent *atomic weight*. Thus the symbols H, Cl, O, etc., not only perform the offices of representing (a) names, (b) single

* Only the atomic weights of the above and a few of the chief metallic elements need be committed to memory; others can be sought out as occasion may require. A complete Table of *combining proportions* of elements, or Atomic Weights, is given at the end of the volume.

volumes, and (c) single atoms, but (d) definite weights of the respective elements. $H=1$, $Cl=35.5$, $O=16$, $I=127$, $N=14$, $K=39$, etc.

LAWS OF CHEMICAL COMBINATION (BY VOLUME).

In 1809 Gay-Lussac showed it to be a fact that when gaseous elements unite with one another in the intimate manner termed chemical, they do not combine in the haphazard proportions (that is, proportions by measure or volume) of a mere mixture, but in constant proportions in the case of any single definite compound, and in simple multiple proportions in cases where two elements form more than one definite compound. He thus proved that the laws respecting the constancy of weight with which elements combine hold good with reference to volume, at all events in those cases in which elements exist in or can be made to assume the gaseous condition. A volume of hydrogen gas and an equal one of chlorine gas give hydrochloric acid gas. Two volumes of hydrogen and one of oxygen give water-vapour or steam. Such volumes or simple multiples are alone the proportions by bulk in which elements combine. If any excess of either gas be mixed and combination attempted, only the stated proportions really combine, the excess remaining unaltered. Further, following Gay-Lussac, on weighing these similar and equal volumes of hydrogen, chlorine, and oxygen, we find that the chlorine is 35.5 times as heavy as hydrogen, and oxygen 16 times as heavy as hydrogen.

In 1811 and 1814 Avogadro and Ampère, reasoning on the fact that all gases are similarly affected by variations of pressure (Boyle, 1662, verified by Mariotte) and temperature (Charles), concluded that all gases must be similarly constituted—similarity in properties always indicating similarity in character or nature; in other words, that if *equal volumes of gases be taken under like conditions*, each will contain the same number of molecules, similar in size and equally distant apart. The deduction is obvious. *The weights of molecules of gaseous elements (that is, of pairs of atoms, and therefore of atoms themselves) must differ to the extent that the weights of equal volumes of those elements differ.* Equal volumes of hydrogen, chlorine, and oxygen, weighing respectively 1, 35.5, and 16, and each of these volumes containing an equal number of molecules, each formed of two atoms, it follows that the relative weights of the atoms will be 1, 35.5, and 16.

It will thus be seen that the weight of the *volume* in which an element combines, and the *actual weight* in which it combines, irrespective of volume, are identical. For instance, we should find by experiment that, as a simple matter of fact, oxygen unites with other elements in proportions of 16 by weight, while hydrogen combines in proportions of 1. Turning, then, to experiments on the volumes in which hydrogen or oxygen combine, and having ascertained those volumes, and then having weighed them, we should find that the oxygen volume weighs 16, while the hydrogen weighs 1. In compounds in which hydrogen were found in proportions of 1 grain, oxygen would be found in proportions of 16 grains. In gaseous compounds in which hydrogen were found in proportions of, say, 27 ounces by measure, oxygen would be found in proportions of 27 ounces by measure; the 27 ounces of hydrogen would be found to weigh 1 grain, and the 27 ounces of oxygen to weigh 16 grains.

Thus the two great facts or laws respecting chemical compounds which Dalton laid down, by ascertaining the exact weights in which bodies combine, Gay-Lussac confirmed by experiments on the exact volumes in which elements combine. Further, Gay-Lussac's experiments and Avogadro's reasoning strongly supported Dalton's theory of atoms.

RECAPITULATION.

What are atomic weights or combining weights? First, they are represented by the smallest portion (relative to 1 part of hydrogen) in which an element migrates from compound to compound. Thus 1 part by weight of hydrogen can be eliminated from 18 similar parts of water by action of certain metals, leaving 1 of hydrogen and 16 of oxygen combined with the metal. From the latter compound 1 more of hydrogen is eliminated by a second experiment with more metal, leaving 16 of oxygen combined with the metal. In these and other well-known reactions 16 parts of oxygen take part in the various operations; 16, therefore, is the probable atomic weight of oxygen; and so with other elements and radicals. Secondly, the weights of the atoms, or the atomic weights of the gaseous elements already studied, must differ from each other to the extent that equal volumes of those elements differ in weight. For equal volumes contain an equal number of molecules equal in size (Avogadro's and Ampère's

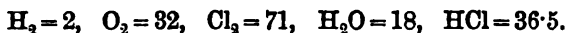
conclusion), and each molecule of an element is composed of two atoms; so that equal volumes of elements contain an equal number of atoms. Now, bulk for bulk, chlorine is thirty-five and a half (35.5) times as heavy as hydrogen; so that the molecule of chlorine must be 35.5 times the weight of the molecule of hydrogen; for molecules are equal in bulk. And as the molecules of chlorine and hydrogen contain two atoms each, the atom of chlorine must be 35.5 times as heavy as that of hydrogen. The actual weight of atoms can never be ascertained; but that is of little consequence if we can only determine, with exactitude, their comparative weights. Comparing, then, all atomic weights (sometimes obscurely termed *equivalents*) with each other, and selecting hydrogen as the standard of comparison (because it is the lightest body known, and therefore, probably, will have the smallest atomic weight), and assigning to it the number 1, we see that the atomic weight of chlorine will be represented by the number 35.5. By parity of reasoning the atomic weight of oxygen is 16; for oxygen is found, by experiment, to be 16 times as heavy as hydrogen. Similarly the atomic weight of nitrogen is found to be 14. The atomic weight of carbon is 12,—not because its vapour has been proved to be 12 times as heavy as hydrogen, for it has never yet been converted into the gaseous state, but because no gaseous compound of carbon which has been analysed has been found to contain in 2 volumes (1 of which, if hydrogen, would weigh 1 part) less than 12 parts of carbon.

By thus weighing equal volumes of gaseous elements, or equal volumes of gaseous compounds of non-volatile elements, and ascertaining by analysis the proportion of the non-volatile element, whose atomic weight is being sought, to the volatile element, whose atomic weight is known, the atomic weights of a large number of the elements have been determined. Some of the elements, however, do not form volatile compounds of any kind; the stated atomic weights of these elements, therefore, are at present simply the proportions by weight in which they combine with or displace elements whose atomic weights have been determined, the proportions being in most cases checked by isomorphic considerations and the relation of the element to other forces, especially heat.* (*Vide infra.*)

* *Isomorphous* bodies (from *isos*, *isos*, equal, and *μορφή*, *morphē*, form) are those which are similar in the shape of their crystals. This identity in crystalline form is so commonly associated with similarity of constitu-

MOLECULAR WEIGHT AND MOLECULAR VOLUME.

The weight of a molecule is simply the sum of the weights of its atoms; thus



Molecular Volume.—If the quantities just mentioned be weighed out (in grains or other weights), or if the molecular weights of any gases or liquids be taken and exposed to similar (high) temperatures and pressures, *they will be found to occupy the same volume.* Conversely, if equal volumes of gases or vapours be measured out, and then the whole weighed, the resulting figures (all referred to 2 of hydrogen as a starting-point or standard) are the molecular weights of the respective substances. Thus a volume of hydrogen (about half a gallon) which, at a temperature of, say, 300° F. or 400° F., and common atmospheric pressure, would weigh 2 grains, would in the case of vapour of water (steam) weigh 18 grains. Hence we are justified in considering, indeed compelled to consider, the molecule of water to contain two atoms of hydrogen (= 2) and one of oxygen (= 16), and its formula to be H_2O (= 18), and not H_4O_2 , in which case its vapour would be twice as heavy as it really is found to be.

Construction of Formulæ.—The composition of hydrochloric acid (HCl), water (H_2O), ammonia gas (NH_3), carbonic acid gas (CO_2), or any other compound, as well as the weight of an element that may be concerned in its formation, cannot be ascertained by actual experiment until the student is far advanced in practical chemistry—until he is able to *analyse* not only *qualitatively*, but, by help of a balance, *quantitatively*. The percentage composition of a chemical substance having been determined by quantitative analysis, its formula is constructed by the aid of the foregoing and other theoretical considerations. The correctness of such formulæ can be verified by expert analysts, but must be taken for granted by learners. The subject will again be referred to in the latter part of this Manual.

tion, that non-crystalline substances resembling each other in structure are often regarded as isomorphous. When one element unites with another in more than one proportion, and its atomic weight is so far uncertain, the isomorphism of either of its compounds with some other compound of known constitution is usually accepted as decisive evidence as to which proportion is atomic. The specific heat of Elements will be treated of subsequently.

QUANTIVALENCE OF ATOMS.

Turning from the *weights* of atoms, their *value* may now be considered, or their *quantivalence* (from *quantitas*, quantity, and *valens*, being worth). The chemical value of atoms in relation to each other may be compared to the exchangeable value of coins. As compared with a penny, a groat is four-valued; as compared with hydrogen, carbon is quadrivalent. Here again hydrogen is conventionally adopted as the standard of comparison. An oxygen atom in its relations to an atom of hydrogen is bivalent (pronounced thus: biv'-a-lent; of double worth, from *bis*, twice, and *valens*); an atom of it will displace two atoms of hydrogen, or combine with the same number; nitrogen is usually trivalent (triv'-a-lent; from *tres*, three, and *valens*); and carbon quad-riv'-a-lent (from *quatuor*, four, or *quater*, four times, and *valens*). Chlorine, iodine, and bromine, as well as potassium, sodium, and silver among the metals, are, like hydrogen, univalent (u-niv'-a-lent; from *unus*, one, and *valens*). Barium, strontium, calcium, magnesium, zinc, cadmium, mercury, and copper, like oxygen, are bivalent. Phosphorus, arsenicum, antimony, and bismuth, like nitrogen, usually exhibit trivalent properties; but the composition of certain compounds of these five elements shows that the several atoms are sometimes quinquivalent (quin-quiv'-a-lent; *quinquies*, five times, and *valens*). Gold and boron are trivalent. Silicon (the characteristic element of flint and sand), tin, aluminium, platinum, and lead resemble carbon in being quadrivalent. Sulphur, chromium, manganese, iron, cobalt, and nickel are sexivalent (sex-iv'-a-lent; from *sex*, six, or *series*, six times, and *valens*), but frequently exert only bivalent, trivalent, or quadrivalent activity. This *quantivalence* (quant-iv'-a-lence; from *quantitas*, quantity, and *valens*), also termed *atomicity* (maximum quantivalence), *dynamicity*, and *equivalence* of elements, may be ascertained at any time on referring to the Table of the Elements at the end of this volume, where Roman numerals, I, II, III, IV, V, VI, are attached to the symbols of each element to indicate atomic univalence, bivalence, trivalence, quadrivalence, quinquivalence, or sexivalence. Dashes (H', O'', N''') similar to those used in accentuating words are often used instead of figures in expressing quantivalence. The quantivalence of elements, as they one after another come under notice, should be carefully committed to memory; for the composition of compounds can often be

thereby predicated with accuracy, and remembered with ease. For instance, the hydrogen compounds of chlorine, Cl' , oxygen, O'' , nitrogen, N''' , and carbon, C'''' , will be respectively $\text{H}'\text{Cl}'$, $\text{H}_2'\text{O}''$, $\text{H}_3'\text{N}'''$, and $\text{H}_4'\text{C}''''$,—one univalent atom, H , balancing or saturating one univalent atom, Cl' ; two univalent atoms, H_2' , and one bivalent atom, O'' , saturating each other; three univalent atoms, H_3' , and one atom having trivalent activity, N''' , saturating each other; and four univalent atoms, H_4' , and one quadrivalent atom, C'''' , saturating each other. Carbonic acid gas, $\text{C}''\text{O}''$, again, is a saturated molecule containing one quadrivalent and two bivalent atoms.

The subject of quantivalence will be further explained after the first six metals have been studied, when abundant illustrations of it will have occurred.

DEFINITIONS.

Chemistry is the study of the chemical force.

The Chemical Force, like other forces, cannot be described, for, like them, it is only known by its effects. It is distinguished from other forces by the facts that (a) it produces an entire change of properties in the bodies on which it is exerted, and (b) that it is exerted only between definite weights and volumes of matter. Like *the force of cohesion*, which is the name given to the attraction which molecules have for each other, and which is great in solids, small in liquids, and apparently absent in gases; and like *the force of adhesion*, which is the name given to the attraction which a mass of molecules has for another mass, *the chemical force* acts only within immeasurable distances; indeed, inasmuch as the chemical force appears to reside in atoms, that is to say is exerted inside a molecule, while all other forces affect entire molecules, the chemical force may be said to be distinguished (c) by being exerted within a smaller distance than that at which any other force is exerted.

An Element is a substance which cannot by any known means be resolved into any simpler form of matter.

An Atom of any element is a particle so small that it undergoes no further subdivision in chemical transformations.

A Molecule is the smallest particle of matter that can exist in a free state.

A mere Mixture of substances is one in which each ingredient retains its properties.

A *Chemical Compound* is one in which definite weights of constituents have undergone an entire change of properties. A "compound" in pharmacy is an intimate mixture of substances, but still only a mixture: it is not a *chemical* compound; the ingredients have not entered into *chemical* union or combination.

Combustion is a variety of chemical combination; a variety in which the chemical union is sufficiently intense to produce heat and, generally, light.

The Law of Diffusion is one under which gases mix with each other at a rate which is in inverse proportion to the square root of their relative weights; that is, irrespective of, and even in spite of, their comparative lightness or heaviness.

A *Chemical Symbol* is a capital letter, or a capital and one small letter. It has four functions, namely—

1. It is short-hand for the *name* of the element.
2. It represents one *atom* of the element.
3. It stands for a constant *weight* of the element—the atomic weight or combining weight.
4. Symbols represent single and equal *volumes* of gaseous elements.

A *Chemical Formula* represents a *molecule* either of an element or of a compound. It has four other functions—

1. It indicates at a glance the *names* of the elements in the molecule.
2. Its symbol, or symbols, together with a small figure attached to the foot of any symbol, show the *number of atoms* in the molecule.
3. It stands for a constant weight of a compound—the molecular weight—the sum of the combining *weights* or of the weights of the atoms in the molecule.
4. It represents *two volumes* of the substance (if volatilizable) in the state of gas or vapour, and the number of volumes of gaseous elements from which two volumes of any gaseous compound were obtained.

A *Chemical Equation* or a *Chemical Diagram* is a collection of formulæ and symbols so placed on paper as to form a picture or illustration of the state of things before and after that *metathesis* (interchange) of atoms of molecules which results in the formation of molecules of new substances.

A *Solid* is a substance the molecules of which are more or less immobile, though probably not in absolute contact.

A *Liquid* is a substance the molecules of which so freely move about each other that it readily assumes and retains the form of any vessel in which it is placed.

A *Gas* is a substance the molecules of which are so far apart that they seem to have lost all attraction for each other, and, indeed, to have acquired the property of repulsion to such an extent that they are only prevented from receding to a still greater extent by the pressure of surrounding matter. Motion is especially characteristic of the molecules of gaseous fluids.

*The Three Laws regulating Chemical Combination
(either by weight or volume).*

First. A definite compound always contains the same elements and the same proportions of those elements—by weight or volume.

Second. When two elements unite in more than one proportion, they do so in simple multiples of that proportion.

Third. The proportions in which two elements unite with a third, are the proportions in which they unite with each other.

Atomic Weights are, first, the proportions in which elements are found to combine with each other by weight. (The figures showing these proportions are purely relative; but all chemists agree to make this relation fixed by giving the number 1 to hydrogen.) Secondly, they are the weights of equal volumes of gaseous elements (relative to one of hydrogen).

Molecular Weights.—These are the weights of equal volumes of gases or vapours, under equal circumstances of temperature and pressure, and relative not to 1 but to 2 of hydrogen. In the case of non-volatile bodies molecular weight is deduced from the observed analogies of the bodies with those whose molecular weight admits of proof.

Quantivalence of Atoms.—The observed power, force, or value for work of an atom—relative to one of hydrogen.

THE LEARNER IS RECOMMENDED TO READ THE FOREGOING PARAGRAPHS ON THE GENERAL PRINCIPLES OF CHEMICAL PHILOSOPHY CAREFULLY ONCE OR TWICE, THEN TO STUDY (EXPERIMENTALLY, IF

POSSIBLE) THE FOLLOWING PAGES, RETURNING TO AND READING OVER THE GENERAL PRINCIPLES FROM TIME TO TIME, UNTIL THEY ARE THOROUGHLY COMPREHENDED.

Minor principles of Chemical Philosophy will be found scattered throughout the following pages.

Students of pure chemistry, especially when fairly well acquainted with chemical facts, will also find the principles of chemistry, including the probable constitution as distinguished from the mere composition of chemical substances, amply set forth in Tilden's "Chemical Philosophy," one of Longmans' series of "Text-books of Science."

QUESTIONS AND EXERCISES.

42. What do you understand by *chemical action*? Give examples.

43. How is the chemical force distinguished from other forces?

44. Adduce evidence that elements exist in compounds; that sulphide of iron, for instance, still contains particles of sulphur and iron, though it possesses properties so different from these elements.

45. Define the term atom.

46. What condition is essential for the manifestation of chemical force?

47. Can an atom exist in an uncombined state? and when are the atoms of an element most potent to enter into chemical combination.

48. What is a molecule?

49. How may the results of chemical reactions be expressed on paper?

50. Enumerate the functions of a symbol.

51. Give the additional functions of a chemical formula.

52. Describe by a diagram or an equation the reaction which ensues when red-hot charcoal is plunged into oxygen gas.

53. Draw diagrams representing the formation of P_2O_5 , SO_2 and FeI_2 respectively.

54. Enumerate the differences in the physical conditions of the molecules in a solid, a liquid, and a gas.

55. State the law of constant proportions.

56. State the law of multiple proportions.
57. State the law of reciprocal proportions.
58. Give illustrations of the above laws.
59. Describe the origin and use of the atomic theory.
60. What do you understand by the atomic weight and the molecular weight of an element?
61. Representing the weight of an atom of hydrogen as 1, what will be the atomic weights of carbon, sulphur, nitrogen, and iodine? Give reasons for considering the stated weights to be correct.
62. In what proportion, by volume, do elements in the gaseous state chemically combine?
63. What relation exists between the combining volumes of elements in the gaseous state and their atomic weights? Give the explanation for this.
64. Is there any difference between the molecular volume of a simple or of a compound gas?
65. Define isomorphism.
66. Explain the value of isomorphism as evidence of atomic weight.
67. What is to be understood by the quantivalence of an element? Give examples of univalent, bivalent, trivalent, and quadrivalent atoms.
68. How may the quantivalence of an element be expressed in its atomic symbol?
69. Give the formulæ of two or three compounds in which the quantivalence of one atom is saturated by the combined quantivalence of others.

The reader is also recommended to question himself, or be questioned, on the "definitions" given on pages 55 to 57.

THE ELEMENTS AND THEIR COMPOUNDS.

Having thus obtained a general idea of the nature of such elements as have special interest for the medical and pharmaceutical student, and which indeed are all with which any student of chemistry should at present occupy his attention, we may pass on to consider in detail the relations of the elements to each other. The elements themselves, in the free

condition, are seldom used in medicine, being nearly always associated—bound together by the chemical force; in this combined condition, therefore, they must be studied. Most compounds met with in the mineral kingdom may be regarded as containing two parts or roots, two radicals:—the one usually metallic, or, to speak more generally, basylous; the other commonly a non-metallic, simple or complex, acidulous radical. In the following pages the basylous radicals, or metals, will be considered first, the acidulous radicals afterwards. Then will follow the chemistry of compounds, many of which have not so simple a constitution as that just indicated. Each radical will be studied from two points of view, the synthetical and the analytical; that is to say, the properties of an element on which the preparation of its compounds depends will be illustrated by descriptions of actual experiments, and thus the principles of chemistry and their applications to medicine and pharmacy be simultaneously learnt; then the reactions by which the element is detected, though combined with other substances, will be performed, and so the student be instructed in qualitative analysis. Synthetical and analytical reactions are, in truth, frequently identical, the object with which they are performed giving them synthetical interest on the one hand, or analytical interest on the other.

A good knowledge of chemistry may be acquired synthetically by preparing considerable quantities of the salts of the different metals, or analytically by going through a course of pure qualitative analysis. But the former plan demands a larger expenditure of time than most students have to spare, while under the latter system pupils generally lose sight of the synthetical interest which attaches to analytical reactions. Hence the more useful system, now offered, of studying each metal, etc., from both points of view, time being economized by the operator preparing only small specimens of compounds.

Chemical synthesis and analysis, thoughtfully and conscientiously followed, without hurry and mere superficial consideration, but, of course, without undue expenditure of time, will insensibly carry the principles of chemistry into the mind, and fix them there indelibly.

POTASSIUM.

Symbol K. Atomic weight 39.

Formula K_2 . Probable molecular weight 78.

Memoranda.—The chief sources of the potassium salts * are the chloride found at Staasfurt, in Prussia, in the form of the mineral *Carnallite* (chloride of potassium 50, chloride of sodium 25, and chloride of magnesium 25, in 100 parts); the nitrate found in soils, especially in warm countries; and the compounds of potassium existing in plants. *Kainit*, a double sulphate of potassium and magnesium, also occurs among the Staasfurt minerals. The vegetable salts of potassium are converted into carbonate (other salts are present) when the wood or other parts are burned to ashes. If the ashes be lixiviated with water, and the solution evaporated to dryness, the residue when fused constitutes *crude potashes*. The residue, calcined on the hearth of a reverberatory furnace till white, gives the product termed *pearlash* or impure carbonate of potassium (the *Potassii Carbonas Impura* of the United States Pharmacopœia). Large quantities of carbonate are thus produced in North America and Russia, and, latterly, from the sugar beetroot marc in France. From the native chloride, and from the carbonate purified "by treating the pearlash with its own weight of distilled water, filtering, and evaporating the solution so formed to dryness, while it is kept briskly agitated" (*Potassæ Carbonas*, B. P., K_2CO_3 , "with about 16 per cent. of water of crystallization"), nearly all other compounds of potassium are made. Exceptions occur in cream of tartar (*Potassæ Tartras Acida*, B. P.), which is, mainly, the purified natural potassium salt of the grape vine, and in nitrate of potassium. Potassium is a constituent of between forty and fifty chemical or Galenical preparations of the British Pharmacopœia.

Carbonate of potassium (*Potassæ Carbonas*, B. P., *Potassii Carbonas*, U. S. P.) is a white crystalline or granular powder, insoluble in alcohol, very soluble in water, rapidly liquefying in the air through absorption of moisture, alkaline and caustic to the taste. It loses all water at a red heat.

Preparation.—Potassium itself is isolated with some diffi-

* The ill-defined term *salt* includes most solid chemical substances, but more especially those which assume a crystalline form.

culty by distilling a mixture of its carbonate and charcoal. It rapidly oxidizes in the air, and hence is always kept below the surface of mineral naphtha, a liquid containing no oxygen. It crystallizes in octohedra.

Quantivalence.—The atom of potassium is univalent, K.

REACTIONS HAVING (a) SYNTHETICAL AND (b) ANALYTICAL INTEREST.

(a) *Synthetical Reactions.*

These are actions utilized in manufacturing preparations of potassium. The word *synthesis* is from σύνθεσις (*sunthēsis*), a putting together, as opposed to analysis from ἀνάλυσις (*analuō*), I resolve.

Hydrate of Potassium. Caustic Potash.

First Synthetical Reaction.—Boil together, for a few minutes, in a test-tube, five or six grains of carbonate of potassium (K_2CO_3) and a like quantity of slaked lime ($Ca2HO$) with a small quantity of water. Set the mixture aside in the test-tube rack till all solid matter has subsided.

This liquid is a solution of caustic potash, or hydrate of potassium (KHO). Made of a prescribed strength, it forms the *Liquor Potassæ*, B.P.

The mixture is known to be boiled long enough when a little of the clear liquid, poured into another test-tube and warmed, gives no effervescence on the addition of an acid (sulphuric, hydrochloric, or acetic)—a test whose mode of action will be explained hereafter.

Best method of expressing decompositions.—This will be easy of comprehension, if what has already been stated concerning symbols and formulæ on pp. 34 to 36 and 41 to 43 has been carefully and thoughtfully considered. The best means of showing on paper the action which occurs when chemical substances attack each other is by the employment either of equations or diagrams, setting forth the formulæ of the molecules concerned in the reaction. In an *equation* the formulæ of the salts used are written on one line, the sign of addition (+) intervening; the sign of equality (=) follows, and then the formulæ of the salts produced, also separated by a plus sign (+). Thus:—



In this reaction (the operation just performed) the metals of (the molecules of) the two salts change places: from K_2CO_3 and $Ca2HO$ there are

produced CaCO_3 and KHO (two molecules, 2KHO); from carbonate of potassium and hydrate of calcium there result carbonate of calcium (the insoluble portion) and hydrate of potassium (in solution).*

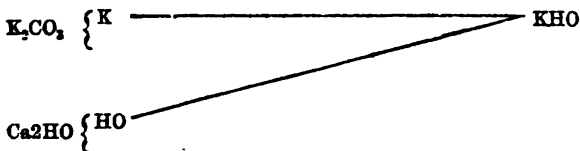
In constructing a *diagram* or pictorial illustration of a chemical reaction (the reaction, for instance, just described), *first* the formulæ of the salts used are written under each other on the left side of the leaf of a note-book, thus:—



Such formulæ are, in this Manual, always given with the description of the reaction. *Secondly*, on the right is then written the formula of the chief substance produced, thus:—



Thirdly, the formation of this chief body under consideration—that is to say, both the origin of its elements and their destination—is traced out by the help of brackets (which show the source of the elements) and converging lines (which suggest the approach and final union of the elements), thus:—



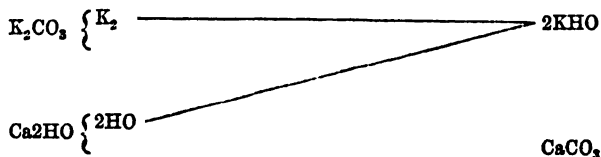
At this stage (at other stages, perhaps, in other reactions) the reader's own intelligent power of thought and reflection must come into exercise. He must reason somewhat as follows:—"I am converting, and entirely converting, a quantity of carbonate of potassium into hydrate of potas-

* If the student is already accustomed to the use of ordinary equations, he may pass on to note 1, on page 65. If not, the author would strongly recommend the temporary employment of diagrams for expressing chemical changes. Indeed, the occasional, if not the regular, use of graphic equations or diagrams is of advantage to all students. For while equation or diagram equally well records the formulæ of the salts concerned in the whole reaction, the diagram alone suggests the *mode* in which its writer believes the respective atoms to change their positions. In the paragraphs succeeding the above, detailed explanations are given respecting the use and construction of diagrams.

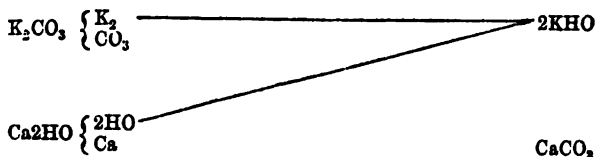
sium. A molecule, the smallest quantity I can picture on paper, of the carbonate of potassium (K_2CO_3) contains, I am told, two atoms of potassium (K_2), and a molecule of the hydrate (KHO) one atom (K). Therefore—therefore—each molecule of the carbonate (K_2CO_3) will furnish two molecules of the hydrate ($2KHO$). Moreover I notice that in the formula of a molecule of the hydrate of calcium (slaked lime) I employ, there are 2 of the HO (that is, $2HO$); and this fact confirms me in the deduction that one molecule of the carbonate affords two molecules of the hydrate." The pupil will then amend his diagram, thus :—



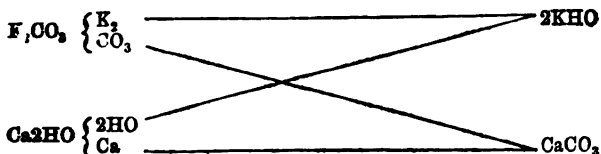
Fourthly, the question as to what becomes of the other elements must be cleared up. Indeed, when the reader remembers that he is studying this reaction for the aid it affords him in learning chemistry, and not because he is desirous of manufacturing caustic potash, he will see that this latter part of the reaction is quite as important as the former. To complete the diagram, then, he must first know what other compound is produced, and its formula. The context of his Manual will generally afford this information, or, after a little experience is acquired, analogies or his own knowledge will suggest correct formulæ. In this case carbonate of calcium is produced, $CaCO_3$. (This product is, in fact, precipitated chalk, together with any excess of slaked lime and any natural impurities in the slaked lime. *Pure* "precipitated chalk" is made by an analogous reaction described subsequently.) The source of the elements of the carbonate of calcium and, finally, their union must be indicated just as the source and mode of formation of the potash were indicated. That is to say, after the formula of this second substance produced ($CaCO_3$) is written on the right of the diagram, thus :—



the source of its elements is shown by writing the symbols for those elements on the right of the bracket attached to the formula containing the symbols of the elements, thus :—



Lines converging from the symbols of the elements and uniting at the formula of the substance are then drawn, to suggest approach of the atoms of the elements and their union to form a molecule of the compound. The diagram will now be complete, thus:—



The formation of a third product or a fourth product would be indicated in a similar manner.

Note 1.—It will be seen that the chief data required in making either equations or diagrammatic notes of decompositions are the symbolic formulae of the various compounds employed and produced. These formulae are, in this Manual, given whenever necessary. Chemists obtain them in the first instance by help of quantitative analysis. By the same means is obtained a check on the probabilities respecting the relative number of molecules concerned in a reaction.

Note 2.—While an equation or a diagram is an attempt to picture the reaction which ensues when molecules of different substances act upon one another, it necessarily only represents two or a minimum number of the molecules. The student will, of course, understand that what is true of these two or three molecules is true of the thousands or millions of molecules forming the mass or whole quantity of material on which he experiments.

Note on Nomenclature.—*Hydrates* are bodies indirectly or directly derived from water by one half of its hydrogen becoming displaced by an equivalent quantity of another radical. Thus, a piece of potassium thrown on to water (HHO) instantly liberates hydrogen, hydrate of potassium (KHO) being formed. The temperature produced at the same time is sufficiently high to cause ignition of the hydrogen, which burns with a purple flame (owing to the presence of a little vapour of potassium), while the hydrate of potassium remains dissolved in the bulk of the water. This radical or root or group of elements (HO), common to all hydrates, is sometimes termed *hydroxyl*. Water might be termed hydrate of hydrogen or hydroxylide of hydrogen.

Explanation.—With regard to the group of atoms represented by the symbols CO_3 and HO , only a few words need be said here. The former (CO_3) is the grouping (root or radical) found in all carbonates; it is termed the carbonic radical, and is as characteristic of carbonates as potassium (K) is of potassium salts. HO (hydroxyl) is characteristic

of all hydrates. CO_3 is a bivalent root, HO is univalent; hence CO_3 is found united with two univalent atoms, as in carbonate of potassium, K_2CO_3 , or with one bivalent atom, as in carbonate of calcium, CaCO_3 ; and HO is found united in single proportion with univalent atoms, as in hydrate of potassium, KHO , or in double proportion with bivalent atoms, as in hydrate of calcium, Ca_2HO . The quantivalence of a metal has only to be learnt, and the formula of its carbonate and hydrate are ascertained without seeing the formula of either. The formulæ of all other metallic salts are constructed on the same principle. But, beyond committing to memory the formulæ and quantivalence of the various groupings characteristic of carbonates, hydrates, nitrates, sulphates, acetates, etc. (see the following Table), special attention should not at present be devoted to the subject of the constitution of salts, but restricted to what may be called the metallic or basylous side of salts. The formulæ and quantivalence of the chief acidulous groupings referred to, and the symbols and quantivalence of allied elementary bodies are included in the following Table:—

Formulæ and Quantivalence of Acidulous Radicals.

All chlorides contain		Cl	} Trivalent radicals.
„ bromides	„	Br	
„ iodides	„	I	
„ cyanides	„	CN	
„ hydrates	„	HO	
„ nitrates	„	NO_3	
„ chlorates	„	ClO_3	} Bivalent radicals.
„ acetates	„	$\text{C}_2\text{H}_3\text{O}_2$	
„ oxides	„	O	
„ sulphides	„	S	
„ sulphites	„	SO_3	
„ sulphates	„	SO_4	
„ carbonates	„	CO_3	} Trivalent radicals.
„ oxalates	„	C_2O_4	
„ tartrates	„	$\text{C}_4\text{H}_4\text{O}_6$	
„ citrates	„	$\text{C}_6\text{H}_5\text{O}_7$	
„ phosphates	„	PO_4	} Trivalent radicals.
„ borates	„	BO_3	

Radicals.—The above elements and compounds are termed *radicals*, each being the common *root* (*radix*) in a series of

salts. Why compound radicals (as NO_3 , SO_3 , PO_4 , etc.) differ in quantivalence cannot well be explained. Their constituent atoms doubtless always exert the same amount of attractive force, nearly but not quite all this force being exerted in retaining the atoms in one group, and the remainder probably determining the quantivalence. Some of the compound radicals are obtainable in the free state, others have yet to be proved capable of isolated existence.*

Pure solution of potash.—Solution of potash generally contains a trace of alumina dissolved from the lime by the hot alkali, but not enough to interfere with the use of the liquid in medicine. If the solution is required for analytical purposes, it may be obtained free from alumina by avoiding the employment of heat, in the manner suggested by Redwood. Half a gallon is made by mixing half a pound of slaked lime with about three pints of water, placing the mixture in a half-gallon bottle (Winchester quart), and adding to it, in small quantities at a time, a solution of half a pound of carbonate of potassium dissolved in the other pint of water, shaking the mixture well for several minutes after each addition. The whole is now set on one side till clear; and then, if a small quantity poured into a test-tube and warmed does not effervesce on the addition of hydrochloric acid, the solution is fit for use. If effervescence (due to carbonic acid gas) occurs, the mixture must be again well shaken. If the lime be good and recently slaked, and the bottle violently shaken once every half-hour, the decomposition will be complete in about ten or twelve hours.

Liquor Potassæ is officially directed to be made as follows:—

Dissolve 1 pound of carbonate of potassium in 1 gallon of water; heat the solution to the boiling point in a clean iron vessel, gradually mix with it 12 ounces of slaked lime, and continue the ebullition for ten minutes with constant stirring. Then remove the vessel from the fire; and when by the subsidence of the insoluble matter the supernatant liquor has become perfectly clear, transfer it by means of a siphon or by decantation to a green-glass bottle furnished with an air-tight stopper, and add distilled water, if necessary, to make it correspond with the tests of *specific gravity* and *neutralizing power*. (These tests and the mode of applying them will be explained in subsequent sections.)

* Some modern authors term these roots *radicles*, a word more usefully expressive of little roots or rootlets.

Solid potash.—Solution of potash evaporated to dryness in a silver or clean iron vessel, and the residue fused and poured into moulds, constitutes *Potassa Caustica*, B. P. It often contains chlorides, detected by nitrate of silver, and sulphates, detected by a barium salt, as described subsequently in connection with hydrochloric and sulphuric acids.

Sulphurated Potash.

Second Synthetical Reaction.—Into a test tube put a few grains of carbonate of potassium previously mixed with half its weight of sulphur. Heat the mixture gradually until it ceases to effervesce. The resulting fused mass poured on a slab and quickly bottled is the *Potassa Sulphurata*, Sulphurated Potash, of the British Pharmacopoeia.

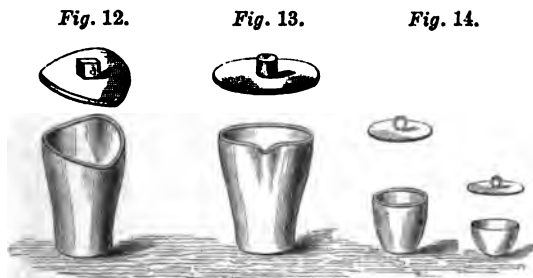


As met with in pharmacy, this salt is not a single definite chemical compound, but a mixture of several; in short, its chemical character is well indicated by its vague name. When fresh, and if carefully prepared with the official proportions of dry ingredients, it is of the colour of liver (whence the old name "liver of sulphur"), and consists, as shown by J. Watts, of the salts mentioned in the foregoing equation, together with a little undecomposed carbonate of potassium, with perhaps higher sulphides of potassium (K_2S_4 and K_2S_5); but, rapidly absorbing oxygen from the air, it soon becomes green and yellow, sulphite (K_2SO_3) and sulphate of potassium (K_2SO_4) are formed, and ultimately a useless mass of a dirty white colour results, consisting of sulphate and hyposulphite, with generally some carbonate of potassium and free sulphur. Moreover, if overheated in manufacture, the hyposulphite $\frac{1}{2}(\text{K}_2\text{S}_2\text{O}_3)$ is decomposed into sulphate $3(\text{K}_2\text{SO}_4)$ and sulphide (K_2S_5) of potassium. Recently made, "about three-fourths of its weight are dissolved by rectified spirit." It is occasionally employed in the form of ointment (*Unguentum Potassæ Sulphurata*, B. P.).

The extremely, indeed most unusually, complicated nature of the decomposition will probably cause failure to any attempt by a student to draw out an equation or a diagram of the reaction without the aid of the

printed equation given above. He may therefore content himself, in this case, by introducing into his note-book a diagram founded directly on the equation and on the numbers of molecules there stated. With this printed equation, and the details of construction of diagrams given in connection with the first synthetical reaction, he will be able to give a diagram of this second synthetical reaction without troubling his reasoning powers, while at the same time he will be familiarizing himself with the more mechanical portions of a diagram.

In preparing large quantities of sulphurated potash, the test-tube is replaced by an earthenware vessel termed a *crucible* (from *crua*, a cross, for originally a cross was impressed upon the melting-pot as used by alchemists and goldsmiths: others derive the word from *crua*, an instrument of torture, the sense here being symbolical).



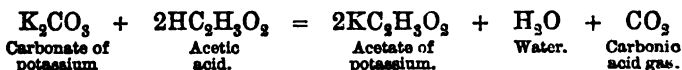
CRUCIBLES OF VARIOUS FORMS.

Heating crucibles.—Crucibles of a few ounces' capacity may be heated in an ordinary grate-fire. Larger ones require a stove with a good draught—that is, a *furnace*. Even the smaller ones are more conveniently and quickly heated in a furnace. Half-ounce or one-ounce experimental porcelain crucibles may be heated in a spirit- or gas-flame; the air-gas flame already described being generally the most suitable.

Acetate of Potassium.

Third Synthetical Reaction.—Place ten, twenty, or more grains of carbonate of potassium in a small dish, and *saturate* (*satur*, full) with acetic acid; that is, add acetic acid so long as effervescence is thereby produced; the resulting liquid is a strong, slightly acid solution of acetate of potassium. Evaporate most of the water in the open dish (*see* figs. 15 & 16,

p. 71), stirring with a glass rod* to promote evolution of vapour; a white salt remains, which fuses on the further application of heat: this is the official Acetate of Potash (*Potassæ Acetas*, B. P.), or Acetate of Potassium (*Potassii Acetas*, U. S. P.) as it is more correctly called. If fused in the open vessel the acetate is liable to become slightly charred and discoloured; this is prevented by transferring the solid residue to a test-tube or florence flask before finally fusing. It forms a white deliquescent foliaceous satiny mass, neutral to test-paper, and wholly soluble in spirit. A ten per cent. solution in water forms the "Solution of Acetate of Potash," B. P.



Explanation of formulæ.—The formula for one molecule of acetic acid (the acetate of hydrogen) is $\text{HC}_2\text{H}_3\text{O}_2$, and one of acetate of potassium $\text{KC}_2\text{H}_3\text{O}_2$. The grouping, $\text{C}_2\text{H}_3\text{O}_2$, is characteristic of all acetates; it is univalent, and may be shortly, though less instructively, written A.

Explanation of process.—When two molecules of acetic acid ($2\text{HC}_2\text{H}_3\text{O}_2$) and one of carbonate of potassium (K_2CO_3) react, two molecules of acetate of potassium ($2\text{KC}_2\text{H}_3\text{O}_2$) and one of carbonic acid (H_2CO_3) are produced, the latter at once splitting up into water (H_2O) and carbonic acid gas (CO_2), as already shown in the equation.

Diagram of the reaction.—The nature of the above operation is indicated by an equation; it (and succeeding reactions) should be expressed in the student's note-book as a *diagram*, and, if possible, without the aid of the equation.

Note.—The above reaction has a general as well as a special synthetical interest. It represents one of the commonest methods of forming salts, namely, the saturation of an acid with a carbonate. Carbonates added to acetic acid yield acetates, to nitric acid nitrates, to sulphuric acid sulphates. Many illustrations of this general process occur in pharmacy.

* Glass rod is usually purchased in the form of long sticks. The pieces may be cut to convenient lengths of from 6 to 12 inches (*vide* p. 5), sharp ends being rounded off by holding in a flame for a few minutes.

Fig. 15.



Fig. 16.

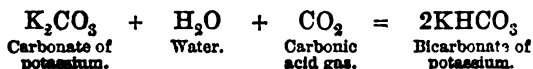


EVAPORATION FROM SMALL AND LARGE BASINS.

Evaporation of water from a liquid is best conducted in wide shallow vessels rather than in narrow deep ones, as the steam can thus quickly diffuse into the air and be rapidly conveyed away; hence a small round-bottomed basin, heated as shown in fig. 15, is far more suitable than a test-tube for such operations. On the manufacturing scale, iron, or iron lined with enamel, or semiporcelain, copper, tinned copper, or solid tin pans are used. Up to 12 or 18 inches diameter, pans, basins, or dishes, made of Wedgwood ware or porcelain composition (fig. 16), may be employed. Small dishes may be supported by retort stands (fig. 15), larger by cylinders (fig. 16) to which the dish is, if less in diameter than the cylinder, adapted by such flat rings or diaphragms as are shown in the figure.

Bicarbonate of Potassium.

Fourth Synthetical Reaction.—Make a *strong* solution of carbonate of potassium by heating in a test-tube a mixture of several grains of the salt with rather less than an equal weight of water. Through the cool solution pass carbonic acid gas, slowly but continuously; after a time a white crystalline precipitate of Acid Carbonate or Bicarbonate of Potassium (KHCO_3), the Bicarbonate of Potash of the Pharmacopœia (*Potassæ Bicarbonas*, B. P., *Potassii Bicarbonas*, U. S. P.), will be formed.



The carbonic acid gas necessary for this operation is to be prepared from marble, though it might be obtained from any carbonate. Thus the previous synthetical reaction could be made available for this purpose, the carbonic gas evolved on the addition of the acetic acid to the carbonate of potassium being conducted into a strong solution of more carbonate of potassium by a glass tube bent and fitted as described when treating of oxygen gas. But motives of economy and convenience induce the use of carbonate of calcium, the form known as marble being always employed. Economy also causes hydrochloric acid to be used in preference to acetic or any other.

Generate the carbonic acid gas by adding common hydrochloric acid, diluted with twice its bulk of water, to a few fragments of marble contained in a test-tube or small flask, and conduct the gas into the solution of carbonate of potassium by a glass tube bent to a convenient angle or angles, and fitted to the test-tube by a cork in the usual way (*see* fig. 10, though no heat is necessary). The tube may be replenished with marble or acid, or both, when the evolution of gas is becoming slow. In working on any larger quantity than a few grains of the carbonate a wide delivery-tube should be employed, or the end of the narrow tube occasionally cleared from any bicarbonate that may have been deposited in it. The more economical official arrangement of the apparatus employed in this process will be described under the corresponding sodium salt (p. 86).

Deposition of the bicarbonate explained.—Bicarbonate of potassium is to a certain extent soluble in water; but as it is less so than the carbonate of potassium, and as a *saturated* solution of the latter has been used, a precipitation of a part of the bicarbonate inevitably occurs. In other words, the quantity of water present is sufficient to keep the carbonate, but insufficient to retain the equivalent quantity of bicarbonate in solution.

Properties.—Prepared on the large scale, bicarbonate of potassium occurs in colourless, non-deliquescent, right rhombic prisms; it has a saline, feebly alkaline, non-corrosive taste.

Effervescing solution of potash.—A solution of 30 grains of bicarbonate of potassium in one pint of water, charged with

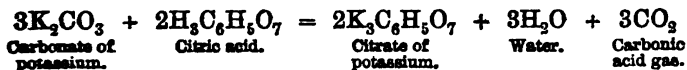
7 times its bulk of carbonic acid gas by pressure, constitutes the ordinary "potash-water," the so-called *Liquor Potassæ Effervescens*, B. P.

Notes on Nomenclature.—The prefix *bi-* in the name "bicarbonate of potassium," serves to recall the fact that to a given amount of potassium this salt contains *twice* as much carbonic radical as the carbonate. The salt is really a "carbonate of potassium and hydrogen" (KHCO_3); it is intermediate between carbonate of potassium (K_2CO_3) and carbonate of hydrogen, or true carbonic acid (H_2CO_3); it is "acid carbonate of potassium" or "hydric potassium carbonate." Hence in constitution it is an acid salt, although not acid to the taste.

Salts whose specific names end in the syllable "*ate*" (carbonate, sulphate, etc.) are in general conventionally so termed when they contain an acidulous radical, or the characteristic elements of an acid, whose name ends in "*ic*," and from which acid they have been or may be formed. Thus the syllable "*ate*," in the words sulphate, nitrate, acetate, carbonate, etc., indicates that the respective salts contain a radical whose name ends in *ic*, the previous syllables, sulph-, nitr-, acet-, carbon-, indicating what that radical is—the sulphuric, nitric, acetic, or carbonic. Occasionally a letter or syllable is dropped from or added to a word to render the name more euphonic; thus the sulphuric radical forms sulphates, not sulphurates.

Citrate of Potassium.

Fifth Synthetical Reaction.—Dissolve a few grains or more of carbonate of potassium in water, and add citric acid ($\text{H}_3\text{C}_6\text{H}_5\text{O}_7$) until it no longer causes effervescence. The resulting liquid is a solution of citrate of potassium ($\text{K}_3\text{C}_6\text{H}_5\text{O}_7$). Evaporated to dryness, in an open dish, a pulverulent or granular residue is obtained, which is the official *Potassæ Citras* (*Potassii Citras*, U. S. P.), a white deliquescent powder.



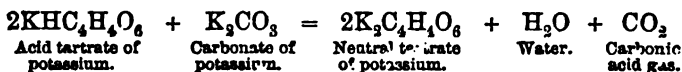
Citrates.—The citric radical or group of elements, which with three atoms of hydrogen forms a molecule of citric acid, and with three of potassium citrate of potassium is a trivalent grouping; hence the three atoms of potassium in a molecule

of the citrate. The full chemistry of citric acid and other citrates will be subsequently described.

Nitrate of potassium (KNO_3) (*Potassæ Nitras*, B. P., *Potassii Nitras*, U. S. P.), and *Sulphate of potassium* (K_2SO_4) (*Potassæ Sulphas*, B. P., *Potassii Sulphas*, U. S. P.), could obviously also be made by saturating nitric acid (HNO_3) and sulphuric acid (H_2SO_4), respectively, by carbonate of potassium. Practically they are not made in that way,—the nitrate occurring, as already stated, in nature, and the sulphate as a by-product in many operations. Both salts will be hereafter alluded to in connection with nitric acid.

Tartrate of Potassium.

Sixth Synthetical Reaction.—Place a few grains of carbonate of potassium in a test-tube with a little water, heat to the boiling-point, and then add acid tartrate of potassium ($\text{KHC}_4\text{H}_4\text{O}_6$ or KHT) till there is no more effervescence; a solution of neutral tartrate of potassium (K_2T) results, the *Potassæ Tartras* of the British Pharmacopœia (*Potassii Tartras*, U. S. P.), the old "Soluble Tartar." Crystals (4- or 6-sided prisms) may be obtained on concentrating the solution by evaporation and setting the hot liquid aside. Larger quantities are made in the same way, 20 of acid tartrate and 9 of carbonate (with 50 of water) being about the proportions necessary for neutrality.



Tartrates.— $\text{C}_4\text{H}_4\text{O}_6$ are the elements characteristic of all tartrates: they form a bivalent grouping; hence the formula of the hydrogen tartrate, or tartaric acid, is $\text{H}_2\text{C}_4\text{H}_4\text{O}_6$; that of the potassium tartrate $\text{K}_2\text{C}_4\text{H}_4\text{O}_6$; of the intermediate salt, the acid potassium tartrate (cream of tartar), $\text{KHC}_4\text{H}_4\text{O}_6$. If the acid tartrate of one metal and the carbonate of another react, a neutral dimetallic tartrate results, as seen in Rochelle Salt ($\text{KNaC}_4\text{H}_4\text{O}_6$), the *Soda Tartarata* of the British Pharmacopœia.

Acid salts (e.g. $\text{KHC}_4\text{H}_4\text{O}_6$), that is, salts intermediate in composition between a normal or neutral salt (e.g. $\text{K}_2\text{C}_4\text{H}_4\text{O}_6$) and an acid (e.g. $\text{H}_2\text{C}_4\text{H}_4\text{O}_6$), will frequently be met with.

All acidulous radicals except those which are univalent may be concerned in the formation of such acid salts.

Iodide of Potassium.

Seventh Synthetical Reaction.—To a solution of potash, heated in a test-tube or flask, or an evaporating-basin, according to quantity, add a small quantity of solid iodine. The deep colour of the iodine disappears entirely. This is due to the formation of the colourless salts, iodide of potassium (KI) and iodate of potassium (KIO₃), which remain dissolved in the liquid. Continue the addition of iodine so long as its colour, after a few minutes' warming and stirring, disappears. When the whole of the potash in the solution of potash has been converted into the salts mentioned, the slight excess of iodine remaining in the liquid will colour it, and thus show that this stage of the operation is completed.

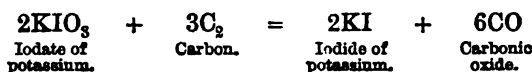


Separation of the iodide from the iodate.—Evaporate the solution to dryness. If each salt were required, the resulting solid mixture might be digested in spirit of wine, which dissolves the iodide, but not the iodate. But the iodide only is used in medicine. Mix the residue, therefore (reserving a grain or two for a subsequent experiment), with about a twelfth of its weight of charcoal, and gently heat in a test-tube or crucible until slight deflagration ensues.* The crucible may be held in a spirit or air-gas flame, or other fire, by tongs. (Scissors-shaped and other "crucible-tongs" are sold

* If, in the operation of heating iodate of potassium with charcoal, excess of the latter be employed, slight incandescence rather than deflagration occurs; if the charcoal be largely in excess, the reduction of the iodate to iodide of potassium is effected without visible deflagration or even incandescence.

Deflagration means violent burning, from *flagratus*, burnt (*flagro*, I burn), and *de*, a prefix augmenting the sense of the word to which it may be attached. Paper thrown into a fire simply burns, nitre deflagrates. Detonate (*detono*) is a precisely similar word, meaning to explode with violent noise.

by all makers of apparatus.) Under these circumstances the iodide remains unaffected; but the iodate loses all its oxygen, and is thus also reduced to the state of iodide.



Treat the mass with a little water, and filter to separate excess of charcoal; a solution of pure iodide of potassium results. The latter may be used as a reagent or it may be evaporated to a small bulk and set aside to crystallize.

This is the process mentioned in the British Pharmacopœia (*Potassii Iodidum*). "Solution of Iodate of Potassium" is also official as a test-liquid.

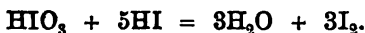
Properties.—Iodide of potassium crystallizes in small cubical crystals, very soluble in water, less so in spirit. One part in ten of water forms "Solution of Iodide of Potassium," B. P. Exposed to air and sunlight, pure iodide of potassium becomes slightly brown, owing to the liberation of iodine. Under these circumstances a little carbonate of potassium is produced by action of the atmospheric carbonic acid, hydriodic acid (HI) is set free, and the latter, attacked by oxygen, yields a trace of water and of free iodine. The ozone in the air (see "Ozone" in Index) may also contribute to the liberation of iodine from such compounds as iodide of potassium.

The addition of charcoal in the above process is simply to facilitate the removal of the oxygen of the iodate of potassium. Iodate of potassium (KIO_3) is analogous in constitution, and in composition, so far as the atoms of oxygen are concerned, to chlorate of potassium (KClO_3), which has already been stated to be more useful than any other salt for the actual preparation of oxygen gas itself. Hence the removal of the oxygen of the iodate might be accomplished by heating the residue without charcoal. In that case the liberated oxygen would be detected on inserting the incandescent extremity of a strip of wood into the mouth of the test-tube in which the mixture of iodide and iodate had been heated. The charcoal, however, burns out the oxygen more quickly, and thus economizes both heat and time.

Note.—The formula of iodide of potassium (KI) shows that the salt contains potassium and iodine in atomic proportions.

A reference to the table of atomic weights at the end of the volume, and a rule-of-three sum, would therefore show what weight of salt is producible from any given weight of iodine.

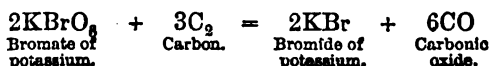
Detection of iodate in iodide of potassium.—Iodate of potassium remaining as an impurity in iodide of potassium may be detected by adding to a solution of the latter salt some weak acid (say, tartaric), shaking, and then adding mucilage of starch; blue "iodide of starch" is formed if a trace of iodate be present, but not otherwise. By the reaction of the added acid and the iodate of potassium iodic acid (HIO_3) is produced, and by reaction of the added acid and the iodide of potassium hydriodic (HI) is produced; neither of these two acids alone attacks starch, but by reaction on each other they give rise to free iodine, which then forms the blue colour. This experiment should be tried on a sample of pure iodide of potassium and on a grain or two of the impure iodide reserved from the previous experiment.



Note on Nomenclature.—The syllable *ide* attached to the syllable *iod* in the name "iodide of potassium," indicates that the *element* iodine is combined with the potassium. An *iodate*, as already explained, is a salt containing the characteristic elements of iodic acid and of all iodic compounds. Salts, one of whose names ends in *ide*, are those which are, or may be, formed from *elements*. The names of salts which are, or may be, formed from compounds include other syllables, *ate* being one (*see* page 73). The only other syllable is *ite*, which is included in the names of salts which are, or may be, formed from acids and radicals whose names end in *ous*: thus hypsulphite of sodium, etc. To recapitulate:—a salt whose name ends in *ate* contains a compound acidulous radical whose name ends in *ic*; a salt whose name ends in *ite* contains a compound acidulous radical whose name ends in *ous*; a salt whose name ends in *ide* contains an element for its acidulous radical. Thus sulphide relates to sulphur, sulphite to the sulphurous radical, sulphate to the sulphuric radical, and so on with all other "ides," "ites," or "ates."

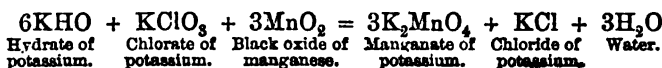
Bromide of Potassium (Potassii Bromidum, B. P.).—This salt is identical in constitution with iodide of potassium, and is

made in exactly the same way, bromine being substituted for iodine. The formula of bromic acid is HBrO_3 . It will be noticed that the following equations are similar in character to those showing the preparation of iodide of potassium :—

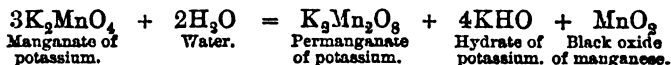


Manganates of Potassium.

Eighth Synthetical Reaction.—Place a fragment of solid caustic potash (KHO), with about the same quantity of chlorate of potassium (KClO_3) and of black oxide of manganese (MnO_2), on a piece of platinum foil.* Hold the foil, by a small pair of forceps or tongs, in the flame of a blowpipe for a few minutes until the fused mixture has become dark green—apparently black. This colour is that of *manganate of potassium* (K_2MnO_4).



Ninth Synthetical Reaction.—*Permanganate of Potassium* ($\text{K}_2\text{Mn}_2\text{O}_8$) (*Potassæ Permanganas*, B. P., *Potassii Permanganas*, U. S. P.), which is purple, is obtained, or rather a solution of it, on placing the foil and its adherent mass in water, and boiling for a short time.



On the large scale, the potash set free in the reaction is neutralized by sulphuric or carbonic acid, and the solution

* The foil may be 1 in. broad by 2 in. long. No ordinary flame will melt the platinum, fused caustic alkalis only slowly corrode it, and very few other chemical substances affect it at all; hence the same piece may be used in experiments over and over again. Most metals form a fusible alloy with platinum, and phosphorus rapidly attacks it; hence such substances, as well as mixtures likely to yield them, should be heated in a small porcelain crucible.

evaporated to the crystallizing point. Further details will be given in connection with manganese.

Solutions of manganate or permanganate of potassium and of sodium so readily yield their oxygen to organic matter, that they are used on the large scale as disinfectants, under the name of "Condy's Disinfecting Fluids."

Synthetical Reactions bringing under consideration the remaining official compounds (namely, bichromate, arsenite, chlorate, cyanide, ferrocyanide, and ferridcyanide of potassium) are deferred at present.

(b) Reactions having Analytical Interest (Tests).

Note.—These are reactions utilized in searching for small quantities of a substance (in the present instance potassium) in a solution. They are best performed in test-tubes or other small vessels. Each should be expressed, in the form of an equation or diagram, in the student's note-book. *All previous or future equations given in this volume should be transferred to the note-book in the form of diagrams, constructed as described on pages 62 to 65, unless the student can with ease construct equations without the aid of the Manual.*

First Analytical Reaction.*—To a solution of any salt of potassium (chloride,† for example) add a few drops of hydro-

* As already indicated, chemical reactions are scarcely analytical or synthetical in themselves, but, rather, performed with an analytical or synthetical object. Indeed, not unfrequently one and the same reaction is both a synthetical and an analytical reaction. Thus this first, so-called, "analytical reaction" is a synthetical reaction if performed with the object of preparing a specimen of the double chloride of platinum and potassium. It is an analytical reaction, or, rather, has analytical interest, if performed with the object of demonstrating the presence of potassium. Chemical reactions in themselves are operations, not so much of analysis (resolution) or synthesis (combination) or of analysis and synthesis conjoined, as of what has sometimes been termed *metathesis* (transposition). Molecules are not torn to atoms in an operation performed with an analytical object, nor are the atoms put together or set together in an operation (perhaps the same operation) performed with a synthetical object; but in both operations the atoms of the molecules undergo metathesis, that is exchange places, or are transposed. In short, chemists use the words "analytical" and "synthetical" in a conventional rather than a strictly etymological sense.

† A few fragments of carbonate of potassium, two or three drops of hydrochloric acid, and a small quantity of water, give a solution of chloride of potassium at once, $K_2CO_3 + 2HCl = 2KCl + H_2O + CO_2$.

chloric acid and of a solution of perchloride of platinum* (Pt Cl_4), and stir the mixture with a glass rod; a yellow granular or slightly crystalline precipitate† slowly forms. (This precipitate is the double chloride of platinum and potassium, and its composition is expressed by the formula $\text{PtCl}_4, 2\text{KCl}$.)

Memoranda.—When the precipitate is long in forming, it is sometimes of an orange-yellow tint. If iodide of potassium happen to be the potassium salt under examination, some iodide of platinum (PtI_4) will also be formed, giving a red colour to the solution, and a larger quantity of the *precipitant* (that is, the precipitating agent) will be required.

Precaution.—Only chloride of potassium forms this characteristic compound: hence, if the potassium salt in the solution is known not to be a chloride, or if its composition is unknown, a few drops of hydrochloric acid must be added, otherwise some of the perchloride of platinum will be utilized for its chlorine only, the platinum being wasted. Thus, if nitrate of potassium (KNO_3) be present, a few drops of hydrochloric acid enable the potassium to assume the form of chloride when the perchloride of platinum is added, nitric acid (HNO_3) being set free.

Explanation.—The precipitate is, practically, insoluble in water. It is for this reason that a very small quantity of any soluble potassium salt (or, rather, of the potassium in that salt) is thrown out of solution by perchloride of platinum.

Note on Nomenclature.—When distinct molecules of salts unite and form a single crystalline compound, the product is termed a *double salt*. The double chloride of potassium and platinum is such a body.

Acid Tartrate of Potassium.

Second Analytical Reaction.—To a solution of any salt of potassium add excess of strong solution of tartaric acid ($\text{H}_2\text{C}_4\text{H}_4\text{O}_6$), and shake or well stir the mixture; a

* Experiments with such expensive reagents as perchloride of platinum are economically performed in watch-glasses, drops of the liquids being operated on.

† By *precipitation* (from *precipitare*, to throw down suddenly) is simply meant the formation of particles of solid in a liquid, no matter whether the solid, the *precipitate*, subsides or floats.

white granular precipitate of acid tartrate of potassium ($\text{KHC}_4\text{H}_4\text{O}_6$) will be formed.

Note.—By “excess” of any test liquid (such as the “solution of tartaric acid” just mentioned) is meant such a quantity as is *probably* rather more than sufficient to convert the whole weight of the compound attacked into the compound produced. Thus, in the present case enough acid must be added to convert the whole of the potassium salt operated on into acid tartrate of potassium. What the weight of salt operated on was must be mentally estimated, roughly, by the operator. It is not necessary in analysing to know the exact weights of salts employed. The analyst must use his judgment, founded on his knowledge of the reaction (as shown by an equation), and of the molecular weights of the substances employed in the reaction, as well as by the rough estimate of the amount of material on which he is experimenting.

Limits of the Test.—Acid tartrate of potassium is soluble in about 180 parts of cold and in 6 parts of boiling water. Hence, in applying the tartaric test for potassium, the solutions must not be hot. Even if cold, no precipitate will be obtained if the solutions are very dilute. This test, therefore, is of far less value than the first mentioned. The acid tartrate of potassium is less soluble in diluted alcohol than in water so that the addition of spirit of wine renders the reaction somewhat more delicate.

Cream of Tartar.—The precipitate is the *Potassæ Tartras Acida* of the British Pharmacopœia (*Potassii Bitartras*, U. S. P.), though the official preparation is not formed in the above manner; on the contrary, the acid is derived from the salt, which, mixed with tartrate of calcium, occurs naturally in the juice of many plants.

Third Analytical Reaction.—The *flame-test*. Dip the looped end of a platinum wire into a solution containing a potassium salt, and introduce the loop into the lower part of a spirit-flame, the flame of a mixture of gas and air, a blowpipe-flame, or other slightly coloured flame. A light violet or lavender tint will be communicated to the flame, an effect highly characteristic of salts of potassium.

Fourth Analytical Fact.—Salts of potassium are not volatile. Place a fragment of carbonate, nitrate, or any other potassium

salt, on a piece of platinum foil, and heat the latter in the flame of a lamp; the salt may fuse to a transparent liquid and flow freely over the foil, water also if present will escape as steam, and black carbon be set free if the salt happen to be a tartrate, citrate, etc.; but the potassium compound itself will not be vaporized. This is a valuable negative property, as will be evident when the analytical reactions of ammonium come under notice.

QUESTIONS AND EXERCISES.

70. Name the sources of potassium.
71. Give the source, formula, and characters of Carbonate of Potassium.
72. Distinguish between synthetical and analytical reactions.
73. How is the official *Liquor Potassæ* prepared?
74. What is the systematic name of Caustic Potash?
75. State the chemical formula of Caustic Potash.
76. Construct an equation or diagram expressive of the reaction between carbonate of potassium and slaked lime.
77. Define a *hydrate*.
78. What group of atoms is characteristic of all carbonates?
79. Define the term *radical*.
80. How is "Sulphurated Potash" made, and of what salts is it a mixture?
81. What is the formula of the acetic radical—the radical of all acetates?
82. Draw a diagram showing the formation of Acetate of Potassium.
83. Give a general process for the conversion of carbonates into other salts.
84. What is the difference between Carbonate and Bicarbonate of Potassium? How is the latter prepared?
85. What is the relation between salts whose specific names end in the syllable "*ate*," and acids ending in "*ic*"?
86. Draw out diagrams or equations descriptive of the formation of Tartrate of Potassium from the Acid Tartrate, and Citrate from the Carbonate of Potassium.
87. Distinguish between a normal and an acid salt.

88. How is Iodide of Potassium made? Illustrate the process either by diagrams or equations.

89. Describe the appearance and chemical properties of iodide of potassium.

89a. Work out a sum showing how much iodide of potassium is producible from 1000 grains of iodine. *Ans.* 1307 grains.

90. Give a method for the detection of iodate in iodide of potassium. Explain the reaction.

91. Has the syllable "*ide*" any general signification in chemical nomenclature?

92. What are the differences between sulphides, sulphites, and sulphates?

93. Mention the chemical relation of Bromide to Iodide of Potassium.

94. Describe the formation of Permanganate of Potassium, giving equations or diagrams.

95. How do manganate and permanganate of potassium act as disinfectants?

96. Enumerate the tests for potassium, explaining by diagrams or equations the various reactions which occur.

SODIUM.

Symbol Na. Atomic weight 23.

Formula Na_2 . Probable molecular weight 46.

Memoranda.—Most of the sodium salts met with in pharmacy are obtained directly from carbonate of sodium, which is now manufactured on an enormous scale from chloride of sodium (common salt, sea-salt, bay-salt, or rock-salt), the natural source of the sodium salts. When pure, salt (*Sodii Chloridum*, B.P.) occurs "in small white crystalline grains, or transparent cubic crystals, free from moisture;" the best varieties commonly contain a little chloride of magnesium, and sometimes other impurities. Besides the direct and indirect use of carbonate of sodium, or carbonate of soda, as it is commonly called, in medicine, it is largely used for household cleansing purposes, under the name of "soda," and in the manufacture of soap. Nitrate of sodium also occurs in nature, but is valuable for its nitric constituents rather than its sodium. Sodium is a constituent of about forty chemical or Galenical preparations of the British Pharmacopoeia.

Sodium is prepared by a process similar to that for potassium, but with less difficulty. It crystallizes in octohedra. Its atom is univalent, Na' .

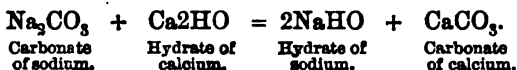
REACTIONS HAVING (a) SYNTHETICAL AND (b) ANALYTICAL INTEREST.

(a) *Reactions having Synthetical Interest.*

Hydrate of Sodium. Caustic Soda.

First Synthetical Reaction.—The formation of solution of hydrate of sodium or caustic soda, NaHO (*Liquor Soda*, B.P.). This operation resembles that of making solution of potash, already described.

The practical student should refer to the remarks made concerning solution of potash, applying them to solution of soda. He may perform the corresponding experiments or omit them, according as he considers he does or does not clearly comprehend all they are designed to teach.



Pure solution of soda, free from any trace of alumina, may be prepared by shaking in a Winchester quart, once every 20 or 30 minutes for 5 or 6 hours, 14 oz. of crystals of carbonate of sodium and 8 oz. of good recently slaked lime. The official *Liquor Soda* is made from 28 ounces of crystals of carbonate of sodium, 12 of slaked lime, and 1 gallon of water, under precisely similar circumstances to those detailed for *Liquor Potassæ* (p. 67). If the solution be evaporated to dryness, and the residue fused and poured into moulds, solid hydrate of sodium (*Soda Caustica*, B. P.) is obtained. Common and cheap caustic soda is now largely employed in various manufactures. This variety is a by-product in the preparation of carbonate of sodium, but, though highly useful as a chemical agent, is too impure for use in medicine.

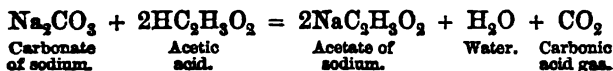
Action of Sodium on Water.—Sodium, like potassium, decomposes water, with production of hydrate of sodium and hydrogen; but unless the sodium is confined to one spot, by placing it on a small floating piece of filter-paper, the action is not sufficiently intense to cause ignition of the escaping

hydrogen. When the latter does ignite, it burns with a yellow flame, due to the presence of a little vapour of sodium.

Second Synthetical Reaction.—The reaction of sulphur and carbonate of sodium at a high temperature resembles that of sulphur and carbonate of potassium; but as the product is not used in medicine the experiment may be omitted. It is mentioned here to draw attention to the close resemblance of the potassium salts to those of sodium.

Acetate of Sodium.

Third Synthetical Reaction.—Add the powder or fragments of carbonate of sodium (Na_2CO_3) to some strong acetic acid in a test-tube or evaporating-basin as long as effervescence occurs, and then evaporate some of the water.* When the solution is cold, crystals of acetate of sodium ($\text{NaC}_2\text{H}_3\text{O}_2 \cdot 3\text{H}_2\text{O}$) (*Sodæ Acetas*, B. P., *Sodii Acetas*, U. S. P.) will be deposited. A ten per cent. solution in distilled water forms the "Solution of Acetate of Soda," B. P.

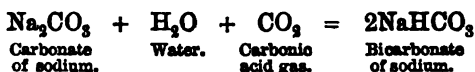


Acetate of sodium effloresces in dry air, and loses all its water of crystallization when gently heated. It supports a temperature of 270° F. or 280° without decomposition, but above 300° soon chars.

Bicarbonate of Sodium.

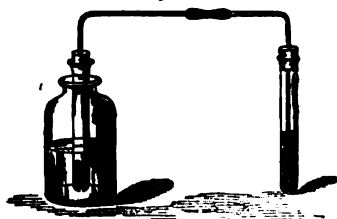
Fourth Synthetical Reaction.—The action of carbonic acid (H_2CO_3), or carbonic acid gas (CO_2) and water (H_2O), on carbonate of sodium (Na_2CO_3). This resembles that of carbonic acid on carbonate of potassium, but is applied in a different manner. The result is bicarbonate of sodium (NaHCO_3) (*Sodæ Bicarbonas*, B. P., *Sodii Bicarbonas*, U. S. P.).

* The "water" alluded to occurs in the acid, which, though commonly termed "acetic acid," is really a solution of that acid in water.



Process.—Heat crystals of carbonate of sodium in a porcelain crucible until no more steam escapes. Rub the product, in a mortar, with two-thirds its weight of more of the crystals, and place the powder in a test-tube or small bottle into which carbonic acid gas may be conveyed by a tube passing through a cork and terminating at the

Fig. 17.



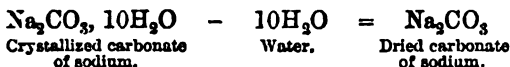
PREPARATION OF BICARBONATE OF SODIUM.

bottom of the vessel. To generate the carbonic acid gas, fill a test-tube having a small hole in the bottom (or a similar piece of glass tubing, of which one end is plugged by a grooved cork) with fragments of marble, insert a cork and delivery-tube, and connect the latter with the similar tube of the vessel containing the carbonate of sodium by a piece of india-rubber tubing. Now plunge the tube of marble into a test glass, or other vessel, containing a mixture of one part hydrochloric acid and 2 parts water, and loosen the cork of the carbonate-of-sodium tube until carbonic acid gas, generated in the marble tube, may be considered to fill the whole arrangement; then replace the cork tightly and set the apparatus aside. As the gas is absorbed by the carbonate of sodium, hydrochloric acid rises into the marble tube and generates fresh gas, which, in its turn, drives back the acid liquid, and thus prevents the production of any more gas until further absorption has occurred. When the salt is wholly converted into bicarbonate (NaHCO_3), it will be found to have become damp through the liberation of water from the crystallized carbonate ($\text{Na}_2\text{CO}_3, 10\text{H}_2\text{O}$). (It would be inconveniently moist, even semi-fluid, if a part of the carbonate had not previously been rendered anhydrous.) To purify the resulting bicarbonate from any carbonate or traces of other salts, add half its bulk of cold

distilled water, set aside for about half an hour, shaking occasionally, drain the undissolved portion, and dry it by exposure on filtering paper.

This is the official process for *Sodæ Bicarbonas*, B. P. The arrangement of apparatus is also that adopted in the *Pharmacopœia* for *Potassæ Bicarbonas*, one part of carbonate dissolved in two and a half parts of water being subjected to the action of the gas, and not the solid carbonate as in the case of the sodium salt.

A crystal of carbonate of sodium is carbonate of sodium plus water; on heating it, more or less of the water is evolved, and anhydrous carbonate of sodium is partially or wholly produced (*Sodæ Carbonas Exsiccata*, B. P., *Sodii Carbonas Exsiccata*, U. S. P.).



Note on Nomenclature.—*Anhydrous* bodies (from α , and $\nu\delta\omega\rho$, $\lambda u d\acute{o}r$, i. e. without water) are compounds from which water has been taken, but whose essential chemical properties are unaltered. Salts containing water are *hydrous* bodies; of these the larger portion are crystalline, and their water is then termed *water of crystallization*. Non-crystalline *hydrous* compounds were formerly spoken of as *hydrated* substances; hydrates are, however, a distinct class of bodies, salts derived from water by one-half of its hydrogen becoming displayed by an equivalent quantity of another radical. *Anhydrides* form still another distinct class of chemical substances; they are derived from acids; in short, they are acids from which, not exactly water as water, but the elements of water have been removed, the essential chemical (acid) properties being thereby greatly altered. (For illustrations, see Index, “Anhydrides.”)

Water of Crystallization.—The water in crystallized carbonate of sodium is in the solid condition, and, like ice and other fusible substances, requires heat for its liquefaction. Many salts (freezing-mixtures) when dissolved in water give a very cold solution. This is because they and their solid water, if they have any, are then, absorbing some heat from surrounding media, converted into liquids. Take away from

water some of its heat, the result is ice. Give to ice (at 32° F.) more heat than it contains already, the result is water (still at 32° F.). (Heat thus taken into a substance without increasing its temperature is said to become *latent*—from *latens*, hiding; it is no longer discoverable by the sense of touch or the thermometer. The term *latent* gives a somewhat incorrect idea, however, of the process; for our knowledge of the extent and readiness with which one form of force is convertible into another renders highly probable the assumption that heat is in these cases converted into motion, the latter enabling the molecules of a solid to take up the new positions demanded by their liquid condition.) The only apparent difference between ice and the water in such crystals as carbonate of sodium, is that ice is solid water in the free, and water of crystallization solid water in the combined state. The former can only exist at and below 32° F., the latter may exist at ordinary temperatures. Many salts, however, which unite with little or even no water of crystallization at common temperatures, take up much, according to Guthrie, at very low temperatures, and such salts he has termed *cryohydrates* (*κρυος*, *kruos*, icy cold, frost). On the other hand, all water of crystallization is expelled at high temperatures. In chemical formulæ, the symbols representing water are usually separated by a comma from those representing salts. The crystals of acetate of sodium (of the third reaction) contain water in this loose state of combination—water of crystallization ($\text{NaC}_2\text{H}_3\text{O}_2, 3\text{H}_2\text{O}$).

“*Soda-water*.”—A solution of bicarbonate of sodium in water charged with carbonic acid gas under pressure constitutes the official *Liquor Sodæ Effervescens*, B. P., and, like the “potash-water” of the shops, is a true medicine, an antacid. Ordinary “soda-water,” however, is in many cases simply a solution of carbonic acid gas in water, and would be more appropriately termed “aërated water”: any medicinal effect it may possess is due to the sedative influence of its carbonic acid gas on the coats of the stomach. At common temperatures water dissolves about its own volume of carbonic acid gas, both being under the same pressure. One pint of the official soda-water contains 30 grains of bicarbonate of sodium and a pint of carbonic acid gas; but the solution is under a pressure of seven atmospheres, so that seven pints of the gas at ordinary atmospheric pressure are required for the quantity mentioned.

Solubility of gases in water.—Whatever the weight and volume of a gas dissolved by a liquid at ordinary atmospheric pressure, that weight is doubled by double pressure, the two volumes of gas thereby being reduced to one; trebled at treble pressure, the three volumes of gas being reduced to one; quadrupled at quadruple pressure, the four volumes of gas being reduced to one, and so on. This is a general law (Henry and Dalton), regarding the solubility of gases in liquids under given temperatures. An average bottle of "soda-water" contains about four times the weight of carbonic acid gas which can exist in it without artificial pressure; so that on removing its cork three times its bulk escapes, its own bulk remaining dissolved.

Nota.—Bicarbonate of sodium may also be medicinally administered in the form of lozenge (*Trochisci Sodæ Bicarbonatis*, B. P.).

Tartrate of Potassium and Sodium.

Fifth Synthetical Reaction.—To some hot strong solution of carbonate of sodium (about three parts), in a test-tube or larger vessel, add acid tartrate of potassium (about four parts) till no more effervescence occurs; when the solution is cold, crystals of the tartrate of potassium and sodium (*Soda Tartrata*, B. P., *Potassii et Sodii Tartras*, U. S. P.), the old *Rochelle Salt*, will be deposited ($\text{KNaC}_4\text{H}_4\text{O}_6 \cdot 4\text{H}_2\text{O}$). The crystals are usually halves of right rhombic prisms.



FORMULÆ OF TARTRATES.

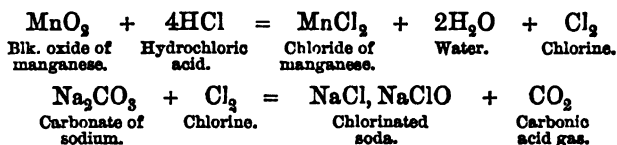
Tartaric acid	HH	$\text{C}_4\text{H}_4\text{O}_6$
Acid tartrate of potassium	KH	$\text{C}_4\text{H}_4\text{O}_6$
Tartrate of potassium and sodium	KNa	$\text{C}_4\text{H}_4\text{O}_6$

Very close analogy will be noticed in the constitution of the molecules of these salts. When the other tartrates come under notice it will be found that they also have a similar constitution.

Hypochlorite of Sodium.

Sixth Synthetical Reaction.—Pass chlorine (*vide* page 18)

into a solution of carbonate of sodium. The result is a bleaching and disinfecting liquid, which, when made of prescribed strength (12 ounces of carbonate in 36 of water, charged by the washed chlorine from 15 fluid ounces of hydrochloric acid and 4 ounces of black oxide of manganese), is the Solution of Chlorinated Soda (*Liquor Sodæ Chloratæ*) of the Pharmacopœia. It is said to contain chloride of sodium (NaCl) and hypochlorite of sodium (NaClO), with some bicarbonate of sodium.



Other Sodium Compounds.

Synthetical Reactions portraying the chemistry of the remaining official compounds (namely, nitrate, sulphate, hyposulphite, borate, arseniate, and valerianate of sodium) are deferred until the several acidulous radicals of these salts have been described.

Phosphate of Sodium.—The preparation and composition of this salt will be most usefully studied after bone-ash, the source of it and other phosphates, has been described. Bone-ash is phosphate of calcium (*see* page 119).

The official Citro-Tartrate (*Sodæ Citro-tartras Effervescens*) is a mixture of bicarbonate of sodium (17 parts), citric acid (6), and tartaric acid (8), heated (to 200° F. or 220°) until the particles aggregate to a granular condition. When required for medicinal use, a dose of the mixture is placed in water; escape of carbonic acid gas at once occurs, and an effervescing liquid results. This substance may be regarded as the official representative of the popular "Effervescing Citrate of Magnesia," so-called, which will be further noticed in connection with the salts of magnesium (page 130).

Soda Powders (*Pulveres Effervescentes*, U. S. P.) are formed of 30 grains of bicarbonate of sodium and 25 of tartaric acid, wrapped separately in papers of different colours. When mixed with water carbonic acid gas escapes, and tartrate of sodium ($\text{Na}_2\text{C}_4\text{H}_4\text{O}_6$) results, a little bicarbonate also remaining.

In the manufacture of Carbonate of Sodium from chloride, the source of the sodium is chloride of sodium, and of the carbonic radical carbonate of calcium in the form of limestone. The chloride is first converted into sulphate, the sulphate is then roasted with coal and limestone, and the resulting *black ash* lixiviated. (*Lixivia*, from *lix*, lye—water impregnated with alkaline salts: hence *lixiviation*, the operation of washing a mixture with the view of dissolving out salts.) The lye, evaporated to dryness, yields crude carbonate of sodium (soda-ash). This process will be further described in connection with Carbonates.

Deliquescence and Efflorescence.—The carbonates of sodium and potassium, chemically closely allied, are readily distinguished physically. Carbonate of potassium quickly absorbs moisture from the air and becomes damp, wet, and finally fluid—it is *deliquescent* (*deliquescent*, melting away). Carbonate of sodium, on the other hand, yields some of its water of crystallization to the air, the crystals becoming white, opaque, and pulverulent—it is *efflorescent* (*efflorescent*, blossoming forth).

Analogy of Sodium salts to Potassium salts.—Other synthetic reactions might be described similar to those given under potassium, and thus citrate, iodide, bromide, iodate, bromate, chlorate, manganate, and permanganate of sodium, and many other salts be formed. But enough has been stated to show how chemically analogous sodium is to potassium. Such analogies will constantly present themselves. In few departments of knowledge is order and method more perceptible; in few is there as much natural law, as much *science*, as in chemistry.

Substitution of Potassium and Sodium salts for each other.—Sodium salts being cheaper than potassium salts, the former may sometimes be economically substituted. That one is employed rather than the other, is often merely a result due to accident or fashion. But it must be borne in mind that in some cases a potassium salt will crystallize more readily than its sodium analogue, or that a sodium salt is stable when the corresponding potassium salt has a tendency to absorb moisture, or one may be more soluble than the other, or the two may have different medicinal effects. For these or similar

reasons, a potassium salt has come to be used in medicine or trade instead of the corresponding sodium salt, and *vice versâ*. Whenever the acidulous portion only is to be utilized, the least expensive salt of the class would nearly always be selected.

(b) *Reactions having Analytical Interest.*

1. *The chief analytical reaction for sodium is the flame-test.* When brought into contact with a flame in the manner described under potassium (page 81), an intensely yellow colour is communicated to the flame by any salt of sodium. This is highly characteristic—indeed, almost too delicate a test; for if the point of the wire be touched by the fingers, enough salt (which is contained in the moisture of the hand) adheres to the wire to communicate a very distinct sodium reaction. These statements should be experimentally verified, the chloride, sulphate, or any other salt of sodium being employed.

2. *Precipitant of Sodium.*—Sodium is the only metal whose common salts are all soluble in water. Hence no ordinary reagent can be added to a solution containing a sodium salt which shall give a precipitate containing the sodium. A neutral or alkaline solution of a sodium salt gives, however, a granular precipitate of antimoniate of sodium ($\text{Na}_2\text{H}_2\text{Sb}_2\text{O}_7, 6\text{H}_2\text{O}$) if well stirred or shaken with a solution of *antimoniate of potassium* ($\text{K}_2\text{H}_2\text{Sb}_2\text{O}_7$). But the reagent precipitates other metals, and is liable to decompose and become useless, and hence is seldom employed.

Antimoniate of potassium is made by adding, gradually, finely powdered metallic antimony to nitrate of potassium fused in a crucible so long as deflagration continues. The resulting mass is boiled with a large quantity of water, the solution filtered and preserved in a well-stoppered bottle; for the carbonic gas in the air is rapidly absorbed by the solution, antimonious acid being deposited.

3. Sodium salts, like those of potassium, are not volatile. Prove this fact by the means described when treating of the effect of heat on potassium salts (p. 82).

QUESTIONS AND EXERCISES.

97. How is the official Solution of Soda prepared? Give a diagram or equation.

98. Explain the action of sodium or potassium on water. What colours do these elements respectively communicate to flame?

99. How much bicarbonate of sodium can be obtained from 2240 pounds of crystallized carbonate of sodium? *Ans.* 1316 lbs.

100. Acetate of Sodium: give formula, process, and equation.

101. Give a diagram showing the formation of Bicarbonate of Sodium.

102. Why is a mixture of dried and undried carbonate of sodium employed in the preparation of the bicarbonate?

103. State the difference between anhydrous and crystallized carbonate of sodium.

104. Define the terms *anhydrous*, *hydrous*, *hydrate*, *anhydride*.

105. What do you understand by *water of crystallization*?

106. What is the nature of the official "*Liquor Sodæ Effervescens*"?

107. How many volumes of gas (reckoned as at ordinary atmospheric pressure) are contained in any given volume of the official "*Soda-water*"?

108. What is the general law regarding the solubility of gases in liquids under pressure?

109. What is the systematic name of *Soda Tartarata*, B. P.? and how is the salt prepared?

110. What is the relation of Rochelle salt to cream of tartar and tartaric acid?

111. Give the mode of preparation and composition of the official Solution of Chlorinated Soda, and express the process by a diagram.

112. How is the granular effervescing Citro-tartrate of Sodium prepared?

113. Define *Deliquescence*, *Efflorescence*, and *Lixiviation*.

114. What is the general relation of potassium salts to those of sodium?

115. How are sodium salts analytically distinguished from those of potassium?

AMMONIUM.

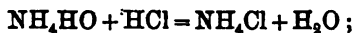
Symbol NH_4 , or Am. Atomic weight 18.

Memoranda.—The elements nitrogen and hydrogen, in the proportion of one atom to four (NH_4), are those characteristic of all the compounds about to be studied, just as potassium (K) and sodium (Na) are the characteristic elements of the potassium and sodium compounds. Ammonium is a univalent nucleus, root, or radical, like potassium or sodium; and the ammonium compounds closely resemble those of potassium or sodium. In short, if, for an instant, potassium or sodium be imagined to be compounds, the analogy between these three series of salts is complete. Ammonium is said to have been isolated, by Weyl, as an unstable dark blue liquid, possessing a metallic lustre.

Source.—The source of nearly all the ammoniacal salts met with in commerce is ammonia gas (NH_3), obtained in distilling coals in the manufacture of ordinary illuminating gas. It is doubtless derived from the nitrogen of the plants from which the coal has been produced. It is possible, however, to produce ammonia from its elements. Thus, coal-dust, air, and vapour of water, all at a red heat, yield, according to Rickman and Thompson, gaseous ammonia. Salt added to the mixture prevents the further combustion of the formed ammonia, and chloride of ammonium sublimes.

Ammonia.—When this gas (NH_3) comes into contact with water (H_2O) in the process of washing and cooling coal-gas, hydrate of ammonium (NH_4HO , or AmHO) is believed to be formed, the analogue of hydrate of potassium (KHO) or sodium (NaHO). The grounds for this belief are the observed analogy of the well-known ammoniacal salts to those of potassium and sodium, the similarity of action of solutions of potash, soda, and ammonia on salts of most metals, and the existence of crystals of an analogous sulphur salt (NH_4HS).

Chloride of Ammonium.—The “ammoniacal liquor” of the gas-works is usually neutralized by hydrochloric acid, by which crude *chloride of ammonium* (sal-ammoniac) is produced,



and from this salt, purified, the others used in pharmacy are directly or indirectly made. Chloride of Ammonium (*Ammonii Chloridum*, B. P.) occurs “in colourless, inodorous, translu-

cent fibrous masses, tough, and difficult to powder, soluble in water [1 in 10 is the "Solution of Chloride of Ammonium," B. P.] and in rectified spirit."

Chloride of ammonium generally contains slight traces of oxychloride of iron, tarry matter, and possibly chlorides of compound ammoniums (*vide* "Artificial Alkaloids" in Index).

Sulphate of Ammonium, $(\text{NH}_4)_2\text{SO}_4$, results when "ammoniacal liquor" is neutralized by sulphuric acid. It is largely used as a constituent of artificial manure; and when purified by recrystallization, is employed in pharmacy (*Ammonii Sulphas*, U. S. P.).

Volcanic Ammonia.—A very pure form of ammonia is that met with in volcanic districts, and obtained as a by-product in the manufacture of borax; the crude boracic acid as imported contains about 10 per cent. of ammonium salts, chiefly sulphate, and double sulphates of ammonium with magnesium, sodium, and manganese (*Howard*).

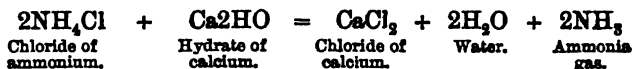
REACTIONS HAVING (a) GENERAL, (b) SYNTHETICAL, AND (c)
ANALYTICAL INTEREST.

Ammonium Amalgam (?)

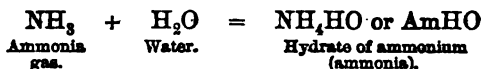
(a) *General Reaction*.—To forty or fifty grains of dry mercury in a dry test-tube, add one or two small pieces of sodium (freed from adhering naphtha by gentle pressure with a piece of filter-paper), and amalgamate by gently warming the tube. To this amalgam, when cold, add some fragments of chloride of ammonium and a strong solution of the same salt. The sodium amalgam soon begins to swell and rapidly increase in bulk, probably overflowing the tube. The light spongy mass produced is the so-called ammonium amalgam, and the reaction is usually adduced as evidence of the existence of ammonium. The sodium of the amalgam unites with the chlorine of the chloride of ammonium, while the ammonium is supposed to form an amalgam with the mercury. As soon as formed the amalgam gives off hydrogen and ammonia gases; this decomposition is nearly complete after some minutes, and impure mercury remains.

(b) *Reactions having Synthetical Interest.***Hydrate of Ammonium. Ammonia.**

First Synthetical Reaction.—Heat a few grains of sal-ammoniac with about an equal weight of hydrate of calcium (slaked lime) damped with a little water in a test-tube; ammonia gas is given off, and may be recognised by its well-known odour. It is very soluble in water. Pass a delivery-tube, fitted to the test-tube as described for the preparation of oxygen and hydrogen, into a second test-tube, at the bottom of which is a little water; again heat, the end of the delivery-tube being only just beneath the surface of the water (or, possibly all the water might rush back into the generating-tube, water absorbing ammonia gas with great avidity); solution of ammonia (*Liquor Ammoniac*, B. P., or *Liquor Ammoniac Fortior*, B. P.) will be thus formed.



Ammonia gas is composed of one atom of nitrogen with three atoms of hydrogen; its formula is NH_3 ; two volumes of it contain one volume of nitrogen combined with three similar volumes of hydrogen. Its constituents have therefore in combining suffered condensation to one-half their normal bulk. Its conversion into hydrate of ammonium may be thus shown:—



Solutions of Ammonia, prepared by this process on a large scale and in suitable apparatus, are met with in pharmacy—the one (sp. gr. 0.891) containing 32.5 per cent., the other (sp. gr. 0.952) 10 per cent. by weight of ammonia gas, NH_3 , or 66.9 and 20.6 of ammonia, NH_4HO (*Liquor Ammoniac Fortior* and *Liquor Ammoniac*, B. P. One part, by measure, of the former, and two of water form the latter). On the large scale, bottles are so arranged in a series as to condense all the ammonia evolved during the operation (see *British Pharmacopœia*).

Note.—When water is mixed with strong solution of ammonia expansion occurs. Thus, calculating from the stated specific gravities of the official solutions, 3000 parts by measure of the *Liquor Ammoniac* weigh 2877 parts (959×3); whereas 1000 parts by measure of the *Liquor Ammoniac Fortior* (weighing 891 parts) and 2000 measures of water (weighing 2000) weigh 2891 parts. Apparently 1000 volumes of the strong solution of ammonia with 2000 volumes of water yield not only 3000 volumes of mixture, but 14 parts by weight, or 15 volumes to spare, total 3015 volumes—an expansion of about half of one (.5) per cent.

Acetate of Ammonium.

Second Synthetical Reaction.—To acetic acid and water in a test-tube add powdered commercial carbonate (acid carbonate and carbamate) of ammonium until effervescence ceases; the resulting liquid, made of prescribed strength, is the official Solution of Acetate of Ammonium ($\text{NH}_4\text{C}_2\text{H}_3\text{O}_2$) (*Liquor Ammoniac Acetatis*, B. P., *Liquor Ammonii Acetatis*, U. S. P.).



Solution of Acetate of Ammonium could, of course, be just as easily made by reaction of acetic acid and solution of ammonia (AmHO), and indeed it was so directed to be prepared in the British Pharmacopoeia of 1864; but the liquid, owing to absence of dissolved carbonic acid, was considered to be too rapid for general use; hence the return to the older method.

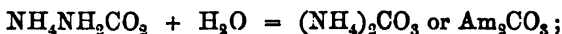
Carbonates of Ammonium.

Commercial carbonate of ammonium is made by heating a mixture of chalk and sal ammoniac; chloride of calcium (CaCl_2) is produced, ammonia gas (NH_3) and water (H_2O) escape, and the ammoniacal carbonate distils or, rather, sublimes* in cakes (*Ammoniac Carbonas*, B. P., *Ammonii Carbonas*, U. S. P.). The best form of apparatus to employ is a retort with a short wide neck and a cool receiver. On the

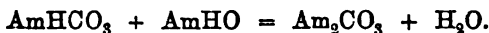
* *Sublimation* (from *sublimis*, high). Vaporization of a solid substance by heat, and its condensation on an upper and cooler part of the vessel or apparatus in which the operation is performed.

large scale the retort is usually iron, and the receiver earthenware or glass; on the small scale glass vessels are employed. The salt is purified by resublimation at a low temperature—150° F. is said to be sufficient.

The salt, the empirical formula of which is $N_4H_{16}C_3O_8$ (B. P.), is probably a mixture of two molecules of acid carbonate of ammonium ($2NH_4HCO_3$) and one of a salt termed carbamate of ammonium ($NH_4NH_2CO_3$). The latter belongs to an important class of salts known as carbamates, but is the only one of interest to the pharmacist. Cold water extracts it from the commercial carbonate of ammonium, leaving the acid carbonate of ammonium undissolved if the amount of liquid used be very small. In water carbamate soon changes into neutral carbonate of ammonium,



so that an aqueous solution of commercial carbonate of ammonium contains both acid carbonate and neutral carbonate of ammonium. If to such a solution some ordinary solution of ammonia be added, a solution of *neutral carbonate of ammonium* is obtained: and this is the common reagent always found on the shelves of the analytical laboratory.



Neutral carbonate of ammonium is the salt formed on adding strong solution of ammonia to the commercial carbonate in preparing a pungent mixture for toilet smelling-bottles; but it is unstable, and on continued exposure to air is reduced to a mass of crystals of the acid carbonate or bicarbonate of ammonium.

According to Divers, the sublimed product of the first distillation of chalk and sal-ammoniac is a mixture of carbamate and carbonate of ammonium, the latter losing some ammonia gas on redistillation, and carbamate with bicarbonate forming the resulting commercial salt. Dr. Divers considers that the salt now met with in trade contains one molecule of acid carbonate to one of carbamate, and not two of the former to one of the latter, as was the case when the analyses were made from which the foregoing formulæ were deduced. Statements by manufacturers and analyses by the author point to the same conclusion.

Sal Volatile (Spiritus Ammoniac Aromaticus, B. P.) is a

spirituous solution of ammonia (AmHO), neutral carbonate of ammonium (Am_2CO_3), and the oils of nutmeg and lemon. Fetid spirit of ammonia (*Spiritus Ammoniae Fœtidus*, B.P.) is an alcoholic solution of the volatile oil of assafoetida mixed with solution of ammonia. "Solution of Carbonate of Ammonia," B.P., is formed by dissolving half an ounce of the salt in ten ounces of water.

Nitrate of Ammonium.

Third Synthetical Reaction.—To some diluted nitric acid add carbonate of ammonium, until, after well stirring, a slight ammoniacal odour remains. The solution contains Nitrate of Ammonium (*Ammoniae Nitras*, B. P., *Ammonii Nitras*, U. S. P.).



Acid carbonate and carbonate
of ammonium.

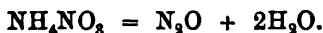
Nitric
acid.

Nitrate of
ammonium.

Water.

Carbonic
acid gas.

From a strong hot solution of nitrate of ammonium crystals may be obtained containing much water ($\text{NH}_4\text{NO}_3 \cdot 12\text{H}_2\text{O}$). On heating these in a dish to about 310°F . the water escapes. The anhydrous salt remaining (NH_4NO_3) may be poured on to an iron plate. On further heating the powdered nitrate, it is resolved into *nitrous oxide* gas (the so-called laughing gas) and water,



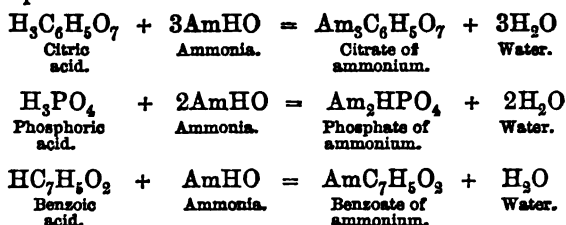
Nitrous oxide is thus prepared for use as an anæsthetic. When required for inhalation, it should be washed from any possible trace of acid or nitric oxide, by being passed through solution of potash and through solution of ferrous sulphate.

Nitrous oxide is slightly soluble in warm water, more so in cold. It supports combustion almost as well as oxygen. By pressure it may be liquefied to a colourless fluid, and by simultaneous cooling solidified.

Citrate, Phosphate, and Benzoate of Ammonium.

Fourth Synthetical Reaction.—To solution of citric acid ($\text{H}_3\text{C}_6\text{H}_5\text{O}_7$ or H_3C_i) add solution of ammonia (AmHO) until the well-stirred liquid smells faintly of ammonia; the product is Solution of Citrate of Ammonium (Am_3C_i) (*Liquor Ammoniae Citratis*, B. P.).

Phosphate of Ammonium (Am_2HPO_4) (*Ammonia Phosphas*, B. P.) and *Benzoate of Ammonium* ($\text{AmC}_7\text{H}_5\text{O}_2$) (*Ammonia Benzoas*, B. P., *Ammonii Benzoas*, U. S. P.) are also made by adding solution of ammonia to phosphoric acid (H_3PO_4) and benzoic acid ($\text{HC}_7\text{H}_5\text{O}_2$) respectively, evaporating (keeping the ammonia in slight excess by adding more of its solution), and setting aside for crystals to form. The official Solution of Acetate of Ammonium, as already stated, could be made in the same way; but when prepared with carbonate of ammonium, the liquid remains charged with carbonic acid, and has a less vivid flavour.



Phosphate of ammonium occurs in transparent colourless prisms, soluble in water, insoluble in spirit; benzoate in crystalline plates, soluble in water and in spirit.

Bromide of Ammonium (*Ammonii Bromidum*, B. P.) will be noticed in connection with Hydrobromic Acid and other Bromides.

Oxalate of Ammonium.

Fifth Synthetical Reaction.—To a nearly boiling solution of 1 part of oxalic acid in about 8 of water add carbonate of ammonium until the liquid is neutral to *test-paper* (see following paragraphs), filter while hot, and set aside for crystals (*Oxalate of Ammonium*, B. P. $(\text{NH}_4)_2\text{C}_2\text{O}_4$, H_2O) to form. The mother-liquor is useful as a reagent in analysis; 1 of the salt in 40 of water constitutes "Solution of Oxalate of Ammonia," B. P.



Neutralization.—Thus far, in reactions, the student has avoided excess of either acid matter on the one hand, or alkali-

line matter on the other, by the rough aid of taste, cessation of effervescence, presence or absence of odour, etc. More delicate aid is afforded by *test-papers*.

Test-papers.—*Litmus* (B. P.) is a blue vegetable pigment, prepared from various species of *Roccella* lichen, exceedingly sensitive to the action of acids, which turn it red. When thus reddened, alkalis (potash, soda, and ammonia) and other soluble hydrates readily turn it blue. The student should here test for himself the delicacy of this action by experiments with paper soaked in solution of litmus and dipped into very dilute solutions of acids, acid salts ($\text{KHC}_4\text{H}_4\text{O}_6$, e.g.), alkalis, and such neutral salts as nitrate of potassium, sulphate of sodium, or chloride of ammonium.

Tincture of Litmus (B. P.).—1 ounce of litmus is macerated for two days in 10 fluid ounces of proof spirit, and the solution poured off from insoluble matter.

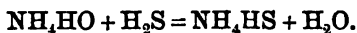
Blue litmus paper (B. P.) is "unsized white paper steeped in tincture of litmus and dried by exposure to the air."

Red litmus paper (B. P.) is "unsized white paper steeped in tincture of litmus which has been previously reddened by the addition of a very minute quantity of sulphuric acid, and dried by exposure to the air."

Turmeric paper, similarly prepared from tincture of turmeric (1 of turmeric root or rhizome to 6 of rectified spirit macerated for seven days), is occasionally useful as a test for alkalis, which turn its yellow to brown; acids do not affect it.

Sulphydrate of Ammonium.

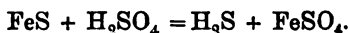
Sixth Synthetical Reaction.—Pass sulphuretted hydrogen gas (H_2S) through a small quantity of solution of ammonia in a test-tube, until a portion of the liquor no longer causes a white precipitate in solution of sulphate of magnesium (Epsom salt); the product is solution of sulphydrate of ammonium (NH_4HS), the "Solution of Sulphide of Ammonium" of the British Pharmacopœia, a most valuable chemical reagent, as will presently be apparent.



"*Solution of Sulphide of Ammonium*," of official strength, is made by passing the gas, prepared in the apparatus described below, into 3 fluid ounces of solution of ammonia (*Liquor Ammoniac*) so long as the gas continues to be absorbed, then

adding 2 more ounces of solution of ammonia, and preserving the solution in a well-stoppered bottle.

Sulphuretted hydrogen is a compound of noxious odour; hence the above operation, and many others, described farther on, in which this gas is indispensable, can only be performed in the open air, or in a *fume-cupboard*—a chamber so contrived that deleterious gases and vapours shall escape into a chimney in connection with the external air. In the above experiment, the small quantity of gas required can be made in a test-tube, after the manner of hydrogen itself. To two or three fragments of sulphide of iron (FeS) add water and then sulphuric acid; the gas is at once evolved, and may be conducted by a tube into the solution of ammonia. Sulphate of iron remains dissolved in the water.



Crystals of sulphhydrate of ammonium (NH_4HS) may be obtained on bringing ammonia gas (NH_3) and sulphuretted hydrogen (H_2S) together at a low temperature. They are soluble in water without decomposition.

Fig. 18.



SULPHURETTED-HYDROGEN APPARATUS.

Sulphuretted-hydrogen Apparatus.—As no heat is necessary in making sulphuretted hydrogen (B. P.), the test-tube of the foregoing operation may be advantageously replaced by a bottle, especially when larger quantities of the gas are required. In analytical operations, the gas should be purified by passing it through water contained in a second bottle.

The most convenient arrangement for experimental use is prepared as follows:—Two common wide-mouth bottles are selected, the one having a capacity of about half a pint, the other a quarter pint; the former may be called the generating-bottle, the latter the wash-bottle. Fit two corks to the bottles. Through each cork bore two holes by a round file or other instrument, of such a size that glass tubing of about the diameter of a quill pen shall fit them tightly. Through one of the holes in the cork of the generating-bottle pass a funnel-tube, so that its extremity may nearly reach the bottom of the bottle. To the other hole adapt a piece of tubing, 6 inches long, and bent in the middle to a right angle. A similar "elbow-tube" is fitted to one of the holes in the cork of the wash-bottle, and another elbow-tube, one arm of which is long enough to reach to near the bottom of the wash-bottle, fitted to the other hole. Removing the corks, two or three ounces of water are now poured into each bottle, an ounce or two of sulphide of iron put into the generating-bottle, and the corks replaced. The elbow-tube of the generating-bottle is now attached by a short piece of india-rubber tubing to the long-armed elbow-tube of the wash-bottle, so that gas coming from the generator may pass through the water in the wash-bottle. The delivery-tube of the wash-bottle is then lengthened by attaching to it, by india-rubber tubing, another piece of glass tubing, several inches in length. The apparatus is now ready for use. Strong sulphuric acid is poured down the funnel-tube in small quantities at a time, until brisk effervescence is established, and more added from time to time as the evolution of gas becomes slow. The gas passes through the tubes into the wash-bottle, where, as it bubbles up through the water, any trace of sulphuric acid, or other matter mechanically carried over, is arrested, and thence the gas flows out at the delivery-tube into any vessel or liquid that may be placed there to receive it. The generator must be occasionally dismantled, and the sulphate of iron washed out.

Luting (lutum, mud). If the corks of the above apparatus are sound, and the tube-holes well made, no escape of gas will

occur. If rough corks have been employed, or the holes are not cylindrical, linseed-meal lute may be rubbed over the defective parts. The lute is prepared by mixing linseed-meal with water to the consistence of dough. A neat appearance may be given to the lute by gently rubbing a well-wetted finger over its surface.

(c) *Reactions having Analytical Interest (Tests).*

First Analytical Reaction.—To a solution of any salt of ammonium (the chloride, for example) in a test-tube, add solution of caustic soda (or solution of potash, or a little slaked lime); ammonia gas is at once evolved, recognised by its well-known odour.



Though ammonium itself cannot be kept in the free state, its compounds are stable. Ammonia is easily expelled from those compounds by action of the stronger alkalies, caustic potash, soda, or lime. As a matter of exercise, the student should here draw out equations in which acetate ($\text{NH}_4\text{C}_2\text{H}_3\text{O}_2$), sulphate (Am_2SO_4), nitrate (NH_4NO_3), or any other ammoniacal salt not already having the odour of ammonia, is supposed to be under examination; also equations representing the use of the other hydrates, potash (KHO) or slaked lime (Ca_2HO).

The odour of ammonia gas is perhaps the best means of recognising its presence; but the following tests are also occasionally useful. Into the test-tube in which the ammonia gas is evolved insert a glass rod moistened with hydrochloric acid (that is, with the solution of hydrochloric acid gas, conventionally termed hydrochloric acid, the *Acidum Hydrochloricum* of the Pharmacopoeia); white fumes of chloride of ammonium will be produced.



Hold a piece of moistened red litmus paper in a tube in which ammonia gas is present; the red colour will be changed to blue.

Second Analytical Reaction.—To a few drops of a solution of an ammonium salt add a drop or two of hydrochloric acid

and a like small quantity of solution of perchloride of platinum (PtCl_4); a yellow crystalline precipitate of the double chloride of platinum and ammonium ($\text{PtCl}_4 \cdot 2\text{NH}_4\text{Cl}$) will be produced, similar in appearance to the corresponding salt of potassium, the remarks concerning which (p. 80) are equally applicable to the precipitate under notice.

Third Analytical Reaction.—To a moderately strong solution of an ammonium salt add a strong solution of tartaric acid, and shake or well stir the mixture; a white granular precipitate of acid tartrate of ammonium will be formed.

For data from which to draw out an equation representing this action, see the remarks and formulæ under the analogous salt of potassium (p. 80).

Fourth Analytical Fact.—Evaporate a few drops of a solution of an ammonium salt to dryness, or place a fragment of a salt in the solid state on a piece of platinum foil, and heat in a flame; the salt is readily volatilised. As already noticed, the salts of potassium and sodium are fixed under these circumstances, a point of difference of which advantage will frequently be taken in analysis. A porcelain crucible may often be advantageously substituted for platinum foil in experiments on volatilization.

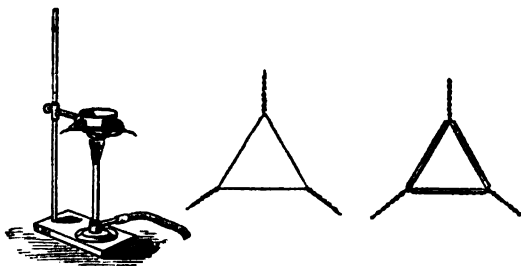
Salts of ammonium with the more complex acidulous radicals seldom volatilize unchanged when heated. The oxalate, when warmed, loses its water of crystallization, and at a higher temperature decomposes, yielding carbonic oxide, carbonic acid gas, ammonia gas, water (the three latter sometimes in combination), and several organic substances. The phosphate yields more or less phosphoric acid as a residue.

A wire triangle may be used in supporting crucibles (fig. 19). It is made by twisting together each pair of ends of three (5- or 6-inch) crossed pieces of wire (fig. 20). A piece of tobacco-pipe stem (about 2 inches) is sometimes placed in the centre of each wire before twisting, the transference of any metallic matter to the sides of the crucible being thus prevented (fig. 21).

Fig. 19.

Fig. 20.

Fig. 21.



TRIANGULAR SUPPORTS FOR CRUCIBLES.

Practical Analysis.

With regard to those experiments which are useful rather as means of detecting the presence of potassium, sodium, and ammonium, than as illustrating the preparation of salts, the student should proceed to apply them to certain solutions of any of the salts of potassium, sodium, and ammonium, with the view of ascertaining which metal is present; that is, proceed to practical analysis.* A little thought will enable him to apply these reactions in the most suitable order and to the best advantage for the contemplated purpose; but the following arrangements are perhaps as good as can be devised:—

DIRECTIONS FOR APPLYING THE FOREGOING ANALYTICAL REACTIONS TO THE ANALYSIS OF AN AQUEOUS SOLUTION OF A SALT OF ONE OF THE METALS, POTASSIUM, SODIUM, AMMONIUM.

Add caustic soda to a small portion of the solution to be examined, and warm the mixture in a test-tube; the odour

* Such solutions are prepared in educational laboratories by a tutor. They should, under other circumstances, be mixed by a friend, as it is not desirable to know previously what is contained in the substance about to be analysed.

The analysis of solutions containing only one salt serves to impress the memory with the characteristic tests for the various metals and other radicals, and familiarize the mind with chemical principles. Medical students seldom have time to go further than this. More thorough analy-

of ammonia gas at once reveals the presence of an ammonium salt.

If ammonium be not present, apply the perchloride-of-platinum test to another portion of the liquid; a yellow precipitate proves the presence of potassium.

(It will be observed that potassium can only be detected in the absence of ammonium, salts of the latter radical giving similar precipitates.)

The flame-test is sufficient for the recognition of sodium.

DIRECTIONS FOR APPLYING THE FOREGOING ANALYTICAL REACTIONS
TO THE ANALYSIS OF AN AQUEOUS SOLUTION OF SALTS OF ONE,
TWO, OR ALL THREE OF THE ALKALI METALS.

Commence by testing a small portion of the solution for an ammonium salt. If it be present, make a memorandum to that effect, and then proceed to get rid of the ammoniacal compound to make way for the detection of potassium; advantage is here taken of the volatility of ammonium salts and the fixity of those of potassium and sodium. Evaporate the original solution to dryness in a small basin, transfer the solid residue to a porcelain crucible, and heat the latter to low redness, or until dense white fumes (of ammoniacal salts) cease to escape (*see* fig. 19). This operation should be conducted in a fume-cupboard, to avoid contamination of the air of the apartment. When the crucible is cold, dissolve out the solid residue with a small quantity of hot water, and test the solution for potassium by the perchloride-of-platinum test, and for sodium by the flame-test.

If ammonium is proved to be absent, the original solution may, of course, be at once tested for potassium and sodium.

Flame-test.—The violet tint imparted to flame by potassium salts may be seen when masked by the intense yellow colour

tical and general chemical knowledge is only acquired by working on such mixtures of bodies as are met with in actual practice, beginning with solutions which may contain any or all of the members of a group. Hence in this Manual two tables of short directions for analysing are given under each group. Pharmaceutical students should follow the second.

due to sodium if the flame be observed through a piece of dark-blue glass, a medium which absorbs the yellow rays of light.

Note on Nomenclature.—The operations of *evaporation* and heating to redness, or *ignition*, are frequently necessary in analysis, and are usually conducted in the above manner. If vegetable or animal matter be also present, carbon is set free, and ignition is accompanied by *carbonization*; the material is said to *char*. When all carbonaceous matter is burnt off, the crucible being slightly inclined and its cover removed to facilitate combustion, and mineral matter, or *ash*, alone remains, the operation of *incineration* has been effected.

Note on the Classification of Elements.—The compounds of potassium, sodium, and ammonium have many analogies. Their carbonates, phosphates, and other common salts are soluble in water. The atoms of the radicals themselves are univalent—that is, displace or are displaced by one atom of hydrogen. In fact, they constitute by their similarity in properties a distinct group or family. All the elements thus naturally fall into classes—a fact that should constantly be borne in mind, and evidence of which should always be sought. It would be impossible for the memory to retain the details of chemistry without a system of classification and leading principles. Classification is also an important feature in the art as well as in the science of chemistry; for without it practical analysis could not be undertaken. The classification adopted in this volume is founded on the quantivalence of the elements and on their analytical relations.

QUESTIONS AND EXERCISES.

116. Why are ammoniacal salts classed with those of potassium and sodium?
117. Mention the sources of the ammonium salts.
118. Describe the appearance and other characters of Chloride of Ammonium.
119. Give the formula of Sulphate of Ammonium.
120. Adduce evidence of the existence of Ammonium.
121. How are the official Solutions of Ammonia prepared? Give diagrams.

122. How is the official Solution of Acetate of Ammonium prepared?

123. What is the composition of commercial Carbonate of Ammonium?

124. Define *sublimation*.

125. What ammoniacal salts are contained in *Spiritus Ammoniac Aromaticus*, B. P., and *Spiritus Ammoniac Fœtidus*, B. P.?

126. Give diagrams or equations illustrating the formation of Citrate, Phosphate, and Benzoate of Ammonium.

127. Give the formula of Oxalate of Ammonium.

128. Show how hydrate of ammonium may be converted into sulphhydrate.

129. Describe the preparation of Sulphuretted Hydrogen gas.

130. Enumerate and explain the tests for ammonium.

131. How is potassium detected in a solution in which ammonium has been found?

132. Give equations illustrating the action of hydrate of sodium on acetate of ammonium; hydrate of potassium on sulphate of ammonium; and hydrate of calcium on nitrate of ammonium.

133. What are the effects of acids and alkalies on litmus and turmeric?

134. Describe the analysis of an aqueous liquid containing salts of potassium, sodium, and ammonium.

135. What meanings are commonly assigned to the terms *evaporation*, *ignition*, *carbonization*, and *incineration*?

136. Write a short article descriptive of the analogies of potassium, sodium, and ammonium, and their compounds.

BARIUM, CALCIUM, MAGNESIUM.

These three elements have many analogies. Their atoms are bivalent.

BARIUM.

Symbol Ba. Atomic weight 137.

The analytical reactions only of this metal are of interest to the general student of pharmacy. The nitrate (Ba_2NO_3) and chloride ($\text{BaCl}_2 \cdot 2\text{H}_2\text{O}$) are the soluble salts in common use in analysis (Chloride of Barium, B. P., and "Solution of Chloride

of Barium," 1 in 10 of water, B. P.) ; and these and others are made by dissolving the native carbonate (BaCO_3), the mineral *witherite*, in acids, or by heating the other common natural compound of barium, the sulphate, *heavy white* or *heavy spar* (CaSO_4), with coal, which yields sulphide of barium (BaS),



and dissolving the sulphide in appropriate acids. When the nitrate is strongly heated it is decomposed, the oxide of barium or *baryta* (BaO) remaining. *Baryta*, on being moistened, assimilates the elements of water with great avidity, and yields hydrate of barium (Ba_2HO). The latter is tolerably soluble, giving *baryta water*; and from this solution crystals of hydrate of barium are obtained on evaporation.

The operations above described may all be performed in test-tubes and small porcelain crucibles heated by the gas-flame. Quantities of 1 oz. to 1 lb. require a coke furnace.

Peroxide of barium (BaO_2) is formed on passing air over *baryta* heated to low redness. By the action of dilute hydrochloric acid it yields solution of *peroxide of hydrogen* (H_2O_2), the old *oxygenated water*.

Quantivalence.—The atom of barium is bivalent, Ba'' .

REACTIONS HAVING ANALYTICAL INTEREST (TESTS).

First Analytical Reaction.—To the solution of any soluble salt of barium (nitrate or chloride, for example) add dilute sulphuric acid; a white precipitate is obtained. Set the test-tube aside for two or three minutes, and when some of the precipitate has fallen to the bottom, pour away most of the supernatant liquid, add strong nitric acid, and boil; the precipitate is insoluble.

The production of a white precipitate by sulphuric acid, insoluble even in hot nitric acid, is highly characteristic of barium. The name of this precipitate is sulphate of barium; its formula is BaSO_4 .

Antidotes.—In cases of poisoning by soluble barium salts, obvious antidotes would be solution of alum or of any sulphates, such as those of magnesium and sodium (Epsom salt, Glauber's salt).

Second Analytical Reaction.—To a barium solution add solution of the yellow chromate of potassium (K_2CrO_4); a pale yellow precipitate ($BaCrO_4$) falls. Add acetic acid to a portion of the chromate of barium; it is insoluble. Add hydrochloric or nitric acid to another portion; it is soluble.

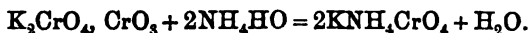
“Neutral Chromate.”—The red chromate (or bichromate) of potassium (K_2CrO_4 , CrO_3) must not be used in this reaction, or the barium will be only imperfectly precipitated; for the red salt gives rise to the formation of free acid, in which chromate of barium is to some extent soluble:—



Yellow chromate is obtained on adding carbonate of potassium, in small quantities at a time, to a hot solution of the red chromate until effervescence ceases; a little more red chromate is then added to insure decomposition of any slight excess of carbonate of potassium.



For analytical purposes solution of a *neutral chromate* is still more readily prepared by simply adding solution of ammonia to solution of red chromate of potassium, until the liquid turns yellow, and, after stirring, smells of ammonia.



Other Analytical Reactions.—To a barium solution add a soluble carbonate (carbonate of ammonia— Am_4CO_3 —will generally be rather more useful than others); a white precipitate of carbonate of barium ($BaCO_3$) results.—To more of the solution add an alkaline phosphate or arseniate (phosphate of sodium— Na_2HPO_4 —is the most common of these chemically analogous salts, but phosphate of ammonium— Am_2HPO_4 —or arseniate— Am_2HAsO_4 —will subsequently have the preference); white phosphate of barium ($BaHPO_4$), insoluble in pure water, but slightly soluble in aqueous solutions of some salts, or arseniate of barium ($BaHAsO_4$), both soluble even in acetic and other weak acids, is precipitated.—To another portion add oxalate of ammonium ($Am_2C_2O_4$); white

oxalate of barium (BaC_2O_4) is precipitated, soluble in the diluted mineral acids, and sparingly so in acetic acid.—The silico-fluoride of barium (BaSiF_6) is insoluble, and falls readily if an equal volume of spirit of wine be added to the solution under examination after the addition of hydrofluosilicic acid (H_2SiF_6).—Barium salts, moistened with hydrochloric acid, impart a greenish colour to flame.

Mem.—Good practice will be found in writing out equations descriptive of each of the foregoing reactions.

QUESTIONS AND EXERCISES.

137. What is the quantivalence of barium?
138. Write down the formulæ of oxide, hydrate, chloride, nitrate, carbonate, and sulphate of barium; and state how these salts are prepared.
139. Describe the preparation of peroxide of hydrogen.
140. Which of the tests for barium are most characteristic? give equations of the reactions.
141. Name the antidote in cases of poisoning by soluble barium salts, and explain its action.

CALCIUM.

Symbol Ca. Atomic weight 40.

Calcium compounds form a large proportion of the crust of our earth. Carbonate of calcium is met with as chalk, marble, limestone, calc-spar, etc.; the sulphate as gypsum or plaster of paris ("Plaster of Paris, native sulphate of calcium— $\text{CaSO}_4, 2\text{H}_2\text{O}$ —deprived of water by heat."—B. P.) and alabaster; the silicate in many minerals; the fluoride of calcium as fluor-spar. The phosphate is also a common mineral. The element itself is only isolated with great difficulty. The atom of calcium is bivalent, Ca".

REACTIONS HAVING SYNTHETICAL INTEREST.

Chloride of Calcium.

First Synthetical Reaction.—To some hydrochloric acid add carbonate of calcium (chalk or, the purer form, white marble, *Marmor Album*, B. P.) (CaCO_3) until effervescence ceases;

filter; solution of chloride of calcium (CaCl_2), the most common soluble salt of calcium, is formed.



This solution contains carbonic acid, and will give a precipitate of carbonate of calcium on the addition of lime-water. It may be obtained quite neutral by well boiling before filtering off the excess of marble. It is a serviceable test-liquid in analytical operations.

Solution of chloride of calcium evaporated to a syrupy consistence readily yields crystals. These are extremely deliquescent. The solution, evaporated to dryness and the white residue strongly heated, gives solid anhydrous chloride of calcium in a porous form. The resulting agglutinated lumps (*Calcii Chloridum*, B. P.) are much used for drying gases, and for freeing certain liquids from water. The salt is also soluble in alcohol. One part of the dried chloride in ten of water constitutes the "Solution of Chloride of Calcium," B. P. Four parts in five of water forms the "Solution (saturated) of Chloride of Calcium," B. P.

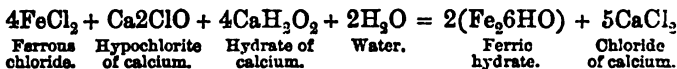
Mem.—The practical student has already met with solution of chloride of calcium as a by-product or secondary product in the preparation of carbonic acid gas.

Marble often contains ferrous carbonate (FeCO_3), which in the above process becomes converted into ferrous chloride, rendering the chloride of calcium impure:—



If absolutely pure chloride of calcium be required, a few drops of the solution should be poured into a test-tube or test-glass, diluted with water, and examined for iron (by adding sulphhydrate of ammonium, which gives a black precipitate with salts of iron), and, if the latter is present, hypochlorite of calcium (in the form of chlorinated lime) and slaked lime be added to the remaining bulk of the liquid, and the whole boiled for a few minutes. Iron (as ferric hydrate) is thus

precipitated; on filtering, a pure solution of chloride of calcium is obtained :—

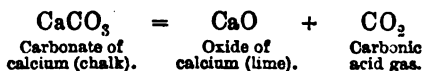


This is the official process, and may be imitated on the small scale after adding a minute piece of iron to a fragment of the marble before dissolving in acid.

The names, formulæ, and reactions of these compounds of iron will be best understood when that metal comes under treatment.

Oxide of Calcium. Quick Lime.

Second Synthetical Reaction.—Place a small piece of chalk in a strong grate-fire or furnace and heat until a trial fragment, chipped off from time to time and cooled, no longer effervesces on the addition of acid; caustic lime, CaO (*Calx*, B. P.), remains.



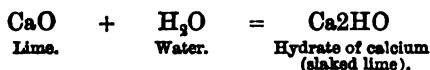
Note.—Etymologically considered, this action is analytical (*ἀναλύω*, *analuō*, I resolve) and not synthetical (*σύνθεσις*, *synthesis*, a putting together); but conventionally it is synthetical, and not analytical; for in this, the usual sense, and the sense in which the words are used throughout the book, synthesis is the application of chemical action with the view of producing something; analysis, the application of chemical action with the view of finding out the composition of a substance. In the etymological view of the matter, there is scarcely an operation performed, either by the analyst or by the manufacturer, but includes both analysis and synthesis—that is, includes interchange or *metathesis*.

Lime-kilns.—On a large scale the above operation is carried on in what are termed *lime-kilns*. (*Kiln*, Saxon *cylŋ*, from *cylene*, a furnace).

Hydrate of Calcium. Slaked Lime.

Slaked lime.—When cold, add to the lime about half its weight of water, and notice the evolution of steam and other evidence of strong action; the product is *slaked lime*, hydrate

of calcium (Ca_2HO) (*Calcis Hydras*, B. P.), with whatever slight natural impurities the lime might contain. The slaking of hard or "stony" lime may be accelerated by using hot water.



Lime-water.—Place the hydrate of calcium in about a hundred times its weight of water; in a short time a saturated solution, known as *lime-water* (*Liquor Calcis*, B. P.), results. It contains about 16 grains of hydrate of calcium (Ca_2HO), equivalent to about 11 or 12 grains of lime (CaO), in one pint.

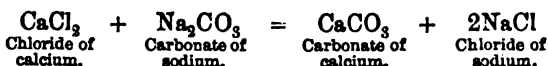
Strong Solution of Lime.—Slaked lime is much more soluble in aqueous solution of sugar than in pure water. The *Liquor Calcis Saccharatus*, B. P., is such a solution, containing 2 ounces of sugar, and 188 grains of hydrate of calcium (Ca_2HO), equivalent to 142 grains of lime (CaO), in 1 pint. It is a more efficient precipitant of hydrates and carbonates than lime-water. The official process is as follows:—Mix 1 ounce of lime and 2 of sugar by trituration in a mortar. Transfer the mixture to a bottle containing 1 pint of water, and, having closed this with a cork, shake it occasionally for a few hours. Finally separate the clear solution with a siphon, and keep it in a stoppered bottle.

Solutions of hydrate of calcium absorb carbonic acid gas on exposure to air, a semi-crystalline precipitate of carbonate being deposited. When the saccharated solution is heated, there is precipitated a compound consisting of three molecules of lime with one of sugar.

Carbonate of Calcium.

Third Synthetical Reaction.—To a solution of chloride of calcium add excess of carbonate of sodium, or about 5 parts of dry chloride to 13 of carbonate; a white precipitate of carbonate of calcium (*Calcis Carbonas Præcipitata*, B. P., *Calcii Carbonas Præcipitata*, U.S.P.) (CaCO_3) results. If the solution of the salts be made hot before admixture, and the whole be set aside for a short time, the particles aggregate to a greater extent than when cold water is used, and the product

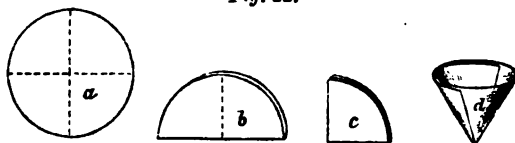
is finely granular or slightly crystalline. The official variety is thus prepared.



Collect and purify this *Precipitated Chalk* by pouring the mixture into a paper cone supported by a funnel, and, when the liquid has passed through the filter, pour water over the precipitate three or four times, until the whole of the chloride of sodium is washed away. This operation is termed *washing a precipitate*. When dried by aid of a *water-bath* (p. 119) or other means, the precipitate is fit for use.

Filtering-paper, or *bibulous paper* (from *bibo*, I drink), is simply good unsized paper made from the best white rags—white blotting-paper, in fact, of unusually good quality. Students' or analysts' filters, on which to collect precipitates, are circular pieces (*a*, fig. 22) of this paper, from three to six inches in diameter, twice folded (*b, c*), and then opened out so as to form a hollow cone (*d*). Square pieces are rounded by scissors *after* twice folding. The cone is supported by a glass or earthenware funnel.

Fig. 22.



CONSTRUCTION OF PAPER FILTERS.

Filters should *always* be cut round, so as to form a cone. If the square piece of paper is folded and used without being so cut or trimmed, an ugly angular filter results, from which it is difficult to wash all "mother-liquor" (the solution of chloride of sodium is the "mother-liquor" in the previous reaction). If a spirituous or other volatile fluid is being passed through such an angular filter, much of the liquid will also be wasted by evaporation from the unnecessarily large surface exposed.

Paper filters of large size are apt to break at the point of the cone. This may be prevented, and the rate of filtration

be much accelerated by supporting the paper cone in a cone of muslin.

Washing-bottle.—Precipitates are best washed by a fine jet of water directed on to the different parts of the filter. A common narrow-necked bottle, of about half-pint capacity (fig. 23), is fitted with a cork; two holes are bored through the cork, the one for a glass tube reaching to the bottom of the bottle within, and externally bent to a slightly acute angle, the other for a tube bent to a slightly obtuse angle, the inner arm terminating just within the bottle. The outer arms may be about 3 inches in length. The extremity of the outer arm continuous with the longer tube should be previously drawn out to a fine capillary opening by holding the original tube, before cutting, in a flame, and, when soft, slowly pulling the halves

Fig. 23.



Fig. 24.



WASHING-BOTTLES.

away from each other until the heated portion is reduced to the thinness of a knitting-needle. The tube is now cut at the thin part by a file, and the sharp edges rounded off by placing in a flame for a second or two. The outer extremity of the shorter tube should also be made smooth in the flame. The apparatus being put together, and the bottle nearly filled with water, air blown through the short tube by the lungs, forces water out in a fine stream at the capillary orifice.

For a hot-water washing-flask (fig. 24) the tubes and cork are fitted to a flask which may be heated. A strip of leather tied round the neck will protect the fingers.

Decantation.—Precipitates may also be washed by allowing them to settle, pouring off the supernatant liquid (fig. 25), agitating with water, again allowing to settle, and so on. This is washing by decantation (*de*, from; *canthus*, an edge). If a stream of liquid flowing from a basin or other vessel exhibits any tendency to run down the outer side of the vessel,

it should be guided by a glass rod placed against the point whence the stream emerges (fig. 26).

If the vessel be too large to handle with convenience, the wash-water may be drawn off by a *siphon*, as shown in miniature in fig. 27. A siphon is a tube of glass, metal, gutta percha, or india-rubber, bent into the form of a V or U, filled with water, and inverted; one end immersed in the wash-water, and the other allowed to hang over the side of the vessel. So long as the outer orifice of the instrument is below the level of any liquid in the vessel, so long will that liquid flow from within outwards.*

Fig. 25.

Fig. 26.

Fig. 27.



DECANTATION.

DECANTATION.

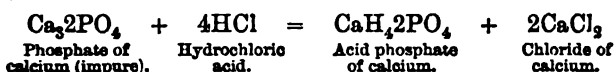
SIPHON IN ACTION.

Prepared carbonate of calcium (*Creta Præparata*, B. P.) is merely washed chalk (*Creta*, B. P.) or *whiting*, only that in Pharmacy fashion demands that the chalk be in little conical lumps, about the size of thimbles, instead of in the larger rolls characteristic of "*whiting*." Wet whiting pushed, portion by portion, through a funnel, and each separately dried, gives the conventional *Creta Præparata*. Its powder is amorphous.

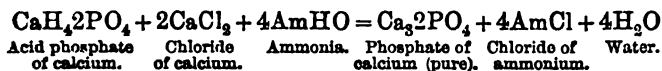
* The nature of the action of a siphon is simple. The column of water in the outer limb is longer, and therefore heavier, than the column of similar area in the inner limb. (The length of the inner limb must be reckoned from the surface of the liquid, the portion below the surface playing no part in the operation.) Being heavier, it naturally falls by gravitation, the liquid in the shorter limb instantly following because pressed upwards by the air. The air, be it observed, exerts a similar amount of pressure on the liquid in the outer limb: in short, atmospheric pressure causes the retention of liquid in the instrument, while gravitation determines the direction of the flow.

Phosphate of Calcium.

Fourth Synthetical Reaction.—Digest bone-ash (bones burnt in an open crucible with free access of air till all animal and carbonaceous matter has been removed,—impure phosphate of calcium—*Os Ustum*, B. P.) with nearly twice its weight of hydrochloric acid (diluted with three or four times its bulk of water) in a test-tube or larger vessel; the phosphate is dissolved.



Dilute with water, filter, boil, and when cold add excess of solution of ammonia; the phosphate of calcium, now pure (*Calcis Phosphas*, B. P., *Calcii Phosphas Precipitata*, U. S. P.), is reprecipitated as a light white amorphous powder. After well washing, the precipitate should be dried over a *water-bath* (see below), or at a temperature not exceeding 212° , to prevent undue aggregation of the particles.



Bone-ash or bone-earth contains small quantities of carbonate and sulphide of calcium. These are decomposed in the above process by the acid, chloride of calcium being formed; on boiling the mixture, carbonic acid gas and sulphuretted hydrogen gas are evolved. Any carbonaceous or siliceous matter, etc., is removed by filtration. In bones the phosphate of calcium is always accompanied by a small quantity of an allied substance, phosphate of magnesium: a trace of fluoride of calcium (CaF_2) is also present.

A *Water-bath* for the evaporation of liquids or for drying moist solids at temperatures below 212°F ., is an iron, tin, or earthenware pan, the mouth of which can be narrowed by iron or tin diaphragms of various sizes, and having orifices adapted to the diameters of basins or plates. (See fig. 16, p. 71.) In the British Pharmacopœia, "when a *water-bath* is directed to be used, it is to be understood that this term refers to an

apparatus by means of which water or its vapour, at a temperature not exceeding 212° , is applied to the outer surface of a vessel containing the substance to be heated, which substance may thus be subjected to a heat near to, but necessarily below, that of 212° . In the *steam-bath*, the vapour of water at a temperature above 212° , but not exceeding 230° , is similarly applied." Evaporation *in vacuo* is performed by simply placing the vessel of liquid over or by the side of a small reservoir of strong sulphuric acid, or other absorbent of moisture, on the plate of an air-pump, covering with a capacious glass hood or "receiver," and exhausting.

Bone-black, or *Animal Charcoal* (*Carbo Animalis*, B. P.), is the residue obtained on subjecting dried bones to a red heat without access of air. The operation may be imitated by heating a few fragments of bone in a covered porcelain crucible in a fume chamber until smoke and vapour cease to be evolved. Purified Animal Charcoal (*Carbo Animalis Purificatus*, B. P.) is obtained by digesting animal charcoal (16 parts) in hydrochloric acid (10 parts) and water (20 parts) in a warm place for a day or two, filtering, thoroughly washing, drying over a water-bath, and igniting the product in a closely covered crucible. The reaction is the same as that just described; that is to say, the acid removes the phosphate of calcium from the carbon of the animal charcoal, by forming soluble acid phosphate and chloride of calcium.

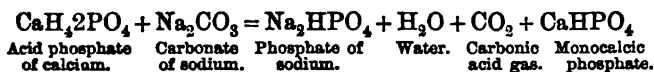
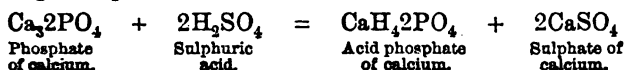
Wood Charcoal (*Carbo Ligni*, B. P.) is wood similarly ignited without access of air.

Decolorizing power of Animal Charcoal.—Animal charcoal, in small fragments, is the material employed in decolorizing solutions of common brown sugar, with the view of producing white lump sugar. Its power, and the nearly equal power of an equivalent quantity of the purified variety, may be demonstrated on solution of litmus or logwood.

Phosphate of Sodium.—Phosphate of calcium is converted into phosphate of sodium (*Sodæ Phosphas*, B. P., *Sodii Phosphas*, U. S. P.) ($\text{Na}_2\text{HPO}_4, 12\text{H}_2\text{O}$) as follows:—Mix, in a mortar, 3 ounces of ground bone-earth with 1 fluid ounce of sulphuric acid; set aside for twenty-four hours to promote

reaction ; mix in about 3 ounces of water, and put in a warm place for two days, a little water being added to make up for that lost by evaporation ; stir in another 3 ounces of water, warm the whole for a short time, filter, and wash the residual sulphate of calcium on the filter to remove adhering acid phosphate of calcium ; concentrate the *filtrate* (the liquid portion), which is a solution of acid phosphate of calcium, to about 3 ounces, filter again if necessary, add solution of (about $4\frac{1}{2}$ ounces of crystals of) carbonate of sodium to the hot filtrate until a precipitate (a phosphate of calcium, CaHPO_4) ceases to form, and the fluid is faintly alkaline ; filter, evaporate, and set aside to crystallize.

Phosphate of sodium occurs "in transparent, colourless, rhombic prisms, terminated by four converging planes ; efflorescent, tasting like common salt." One part in ten of water constitutes "Solution of Phosphate of Soda," B. P. This is the official as well as the ordinary process. The following equations show the two decompositions which occur during the operations :—

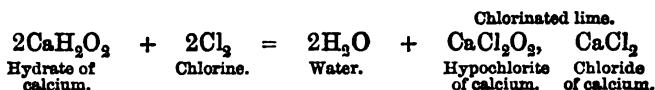


Ordinary phosphate of sodium ($\text{Na}_2\text{HPO}_4, 12\text{H}_2\text{O}$) effloresces rapidly in the air until nearly half its water has escaped, when it has a permanent composition represented by the formula $\text{Na}_2\text{HPO}_4, 7\text{H}_2\text{O}$. Phosphate of sodium has an alkaline reaction. Neutralization by acid results in the removal of half its sodium and formation of the salt $\text{NaH}_2\text{PO}_4, \text{H}_2\text{O}$.

Hypochlorite of Calcium.

Fifth Synthetical Reaction.—Pass chlorine, generated as already described, into damped slaked lime contained in a piece of wide tubing, open at the opposite end to that in which the delivery-tube is fixed. (A test-tube, the bottom of which has been accidentally broken, is very convenient for such opera-

tions.) The product is ordinary *bleaching-powder*, a compound of hypochlorite and chloride of calcium, commonly called *chloride of lime*, the *Calx Chlorata* of the British Pharmacopœia.



Chlorinated lime exposed to air and moisture, as in disinfecting the atmosphere of sick rooms, slowly yields hypochlorous acid (HClO). Free hypochlorous acid soon breaks up into water, chloric acid (HClO_3), and free chlorine. Chloric acid is also unstable, decomposing into oxygen, chlorine, water, and perchloric acid (HClO_4). The small quantity of hypochlorous acid diffused through an apartment when bleaching-powder is exposed thus yields fourteen-fifteenths of its chlorine in the form of chlorine gas—one of the most efficient of known disinfectants.

Constitution of bleaching-powder.—Treated with alcohol, bleaching-powder does not yield its chloride of calcium to the solvent; hence the powder is not a mere mixture of chloride and hypochlorite of calcium: water, also, does not dissolve out first one salt and then the other, but both together, in the molecular proportions of the above formula. On the other hand, when the aqueous solution is cooled, or evaporated *in vacuo*, crystals are obtained which Kingzett has shown to be nearly pure hypochlorite of calcium, the solution containing chloride of calcium. While the former fact indicates that the powder is a compound, and not a mere mixture, the later indicates that it is a feeble compound—an adhesion of molecules of hypochlorite and chloride, as shown in the equation, rather than any more intimate or closer combination of atoms. If it be regarded as a single rather than a double salt, then the following formula may be employed, $\text{Ca} \left\{ \begin{array}{l} \text{Cl} \\ \text{ClO} \end{array} \right.$.

Bleaching-liquor.—Digest chlorinated lime in water, in which the bleaching compound is soluble, filter from the un-

dissolved lime, and test the bleaching powers of the clear liquid by adding a few drops to a decoction of logwood slightly acidulated. One pound of this bleaching-powder, shaken several times during three hours with 1 gallon of water, forms the official Solution of Chlorinated Lime (*Liquor Calcis Chloratae*, B. P.).

Gummate of Calcium.

Gummate of Calcium is the only official calcium salt that remains to be noticed. This compound is, in short, *arabin*, the ordinary Gum-Acacia or Gum-Arabic (*Acaciae Gummi*, B. P.), a substance too well known to need description. A solution of gum-arabic in water (*Mucilago Acaciae*, B. P.) yields a white precipitate of oxalate of calcium on the addition of solution of oxalate of ammonium. Or a piece of gum burnt to an ash in a porcelain crucible yields a calcareous residue, which, dissolved in dilute acids, affords characteristic reactions with any of the following analytical reagents for calcium. In some specimens of gum-arabic a portion of the calcium is displaced by an equivalent quantity of potassium or magnesium. The gummic radical may be precipitated as opaque gelatinous gummate of lead by the addition of solution of oxyacetate of lead (*Liquor Plumbi Subacetatis*, B. P.) to an aqueous solution of gum. These statements may be experimentally verified by the practical student.

Tragacanth (*Tragacantha*, B. P.) is a mixture of soluble arabinoid gum and a variety of calcium gum insoluble in water, termed *bassorin*. With water a gelatinous mucilage is formed (*Mucilago Tragacanthae*, B. P.).

REACTIONS HAVING ANALYTICAL INTEREST (TESTS).

First Analytical Reaction.—Add sulphuric acid, *very* highly diluted, to a calcium solution contained in a test-tube or small test-glass; sulphate of calcium ($\text{CaSO}_4 \cdot 2\text{H}_2\text{O}$) is formed, but is *not* precipitated, it being, unlike sulphate of barium, slightly soluble in water.

Solution of Sulphate of Calcium.—A quarter of an ounce of that (dried) form of sulphate of calcium known as plaster of Paris (CaSO_4) digested in one pint of water for a short time, with occasional shaking, and the mixture filtered, yields the official test-liquid termed "Solution of Sulphate of Lime," B.P. About 400 parts of the solution contain 1 of sulphate of calcium.

Second Analytical Reaction.—Add yellow chromate of potassium (K_2CrO_4) or other neutral chromate (KAmCrO_4) to a calcium solution slightly acidified with acetic acid; chromate of calcium (CaCrO_4) is probably formed, but is *not* precipitated. Barium *is* precipitated by the chromic radical.

These two negative reactions are most valuable in analysis, as every precipitant of calcium is also a precipitant of barium; but the above two reagents are precipitants of barium only. Hence, calcium, which when alone can be readily detected by the following reactions, cannot by any reaction be detected in the presence of barium. But by the sulphuric or chromic test barium is easily removed, and then either of the following reagents will throw down the calcium.

Other Analytical Reactions.—Add carbonate of ammonium, phosphate of sodium, arseniate of ammonium, and oxalate of ammonium to calcium solutions as described under the analytical reactions of barium, and write out descriptive equations. The precipitates correspond in appearance to those of barium; their constitution is also similar, hence their correct formulæ can easily be deduced. Of these precipitants oxalate of ammonium is that most commonly used as a reagent for calcium salts, barium being absent. The oxalate of calcium is insoluble in acetic, but soluble in hydrochloric or nitric acids.—Calcium compounds impart a reddish colour to flame.

QUESTIONS AND EXERCISES.

142. Enumerate some of the common natural compounds of calcium.

143. Explain, by an equation, the action of hydrochloric acid on marble. What official compound results?

144. Why is chloride of calcium used as a desiccator for gases?
145. How would you purify Chloride of Calcium which has been made from ferruginous marble? Give diagrams.
146. Write a few lines on the chemistry of the lime-kiln.
147. In what sense is the conversion of chalk into lime an analytical action?
148. What occurs when lime is "slaked"?
149. To what extent is lime soluble in water (*Liquor Calcis*, B. P.)? to what in Syrup (*Liquor Calcis Saccharatus*, B. P.)?
150. Describe the preparation of the official Precipitated Carbonate of Calcium (*Calcis Carbonas Præcipitata*, B. P.); in what does it differ from Prepared Chalk (*Oreta Præparata*, B. P.)?
151. In what does filtering-paper differ from other kinds of paper?
152. Explain the construction of "a washing-bottle" for cleansing precipitates by water.
153. Define *decantation*.
154. Describe the construction and manner of employment of a siphon.
155. Explain the mode of action of a siphon.
156. What is the difference between *Os Ustum*, B. P., and *Calcis Phosphas*, B. P.?
157. How is "bone-earth" purified for use in medicine?
158. Explain the action of hydrochloric acid on Animal Charcoal in the conversion of *Carbo Animalis*, B. P., into *Carbo Animalis Purificatus*, B. P.
159. What is the chemical difference between *Carbo Animalis*, B. P., and *Carbo Ligni*, B. P.?
160. Give equations showing the conversion of Phosphate of Calcium into Phosphate of Sodium.
161. Write a short article on the manufacture, composition, and uses of "bleaching-powder" (*Calx Chlorata*, B. P.).
162. How may calcium be detected in Gum-Arabic?
163. State the chemical nature of Tragacanth.
164. To what extent is sulphate of calcium soluble in water?
165. Can calcium be precipitated from an aqueous solution containing barium?
166. Barium being absent, what reagents may be used for the detection of calcium? Which is the chief test?

MAGNESIUM.

Symbol Mg. Atomic weight 24.

Source.—Magnesium is abundant in nature as magnesian or mountain limestone, termed *dolomite*, a double carbonate of magnesium and calcium in very common use as a building-stone (*e.g.* the Houses of Parliament and the School of Mines, in London), and *magnesite*, a tolerably pure carbonate of magnesium, though too “stony” for direct use in medicine, even if very finely powdered. Chloride of magnesium and sulphate of magnesium (Epsom salt) also occur in sea-water and the water of many springs. A monohydrous sulphate ($\text{MgSO}_4 \cdot \text{H}_2\text{O}$), termed *kieserit*, occurs near Staasfurt, in Prussia. Metallic magnesium may be obtained from the chloride by the action of sodium. It burns readily in the air, emitting a dazzling light due to the white heat to which the resulting particles of magnesia (MgO) are exposed. The chloride employed as a source of the metal is obtained by dissolving the carbonate in hydrochloric acid, adding some chloride of ammonium, evaporating to dryness, heating the residue in a flask (on the small scale a large test-tube or florence flask) until the chloride of ammonium is all volatilized, and the chloride of magnesium remains as a clear fused liquid. The latter is poured on to a clean earthenware slab. The chloride of ammonium prevents reaction between chloride of magnesium and water in the last stages of the operation, and consequent formation of oxide (or oxychloride) of magnesium and hydrochloric acid gas.

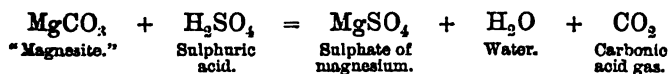
Quantivalence.—The atom of magnesium is bivalent, Mg^2 .

REACTIONS HAVING SYNTHETICAL INTEREST.

Sulphate of Magnesium.

First Synthetical Reaction.—To a few drops of sulphuric acid and a little water in a test-tube, made hot (or to larger quantities in larger vessels), add powdered native carbonate of magnesium, *magnesite*, MgCO_3 , until effervescence ceases, subsequently boiling to aid in the expulsion of the carbonic acid gas. The filtered liquid is a solution of sulphate of magnesium (MgSO_4), crystals of which, *Epsom salt* ($\text{MgSO}_4 \cdot 7\text{H}_2\text{O}$) (*Magnesiæ Sulphas*, B. P., *Magnesiæ Sulphas*, U.S.P.),

may be obtained on evaporating most of the water, and setting the concentrated solution aside to cool. This is an ordinary manufacturing process. Instead of magnesite, *dolomite*, the common magnesian limestone (carbonate of magnesium and of calcium, CaCO_3 , MgCO_3), may be employed, any iron being removed by evaporating the solution (filtered from the sulphate of calcium produced) to dryness, gently igniting to decompose sulphate of iron, dissolving in water, filtering from oxide of iron, and crystallizing. (If neither mineral be at hand, the practical student may use a little of the ordinary manufactured carbonate of pharmacy; for the chemical action is almost identical, and it is the chemistry, and not, just now, the commercial economy of the matter, that he is studying. The manufacturer must, of course, commence with one of the above mineral carbonates furnished by nature, from that make his sulphate, and from the latter, as will be seen directly, make the pure pulverulent carbonate of pharmacy.)



Sulphate of magnesium readily crystallizes in large, colourless, transparent, rhombic prisms; but, from concentrated solutions, the crystals are deposited in short thin needles, a form more convenient for manipulation, solution, and general use in medicine.

Iron may be detected in sulphate of magnesium by adding the common alkaline solution of chlorinated lime or chlorinated soda to some aqueous solution of the salt, brown hydrate of iron (Fe_3HO) being precipitated. Sulphydrate of ammonium will also give a black precipitate if iron be present.

Carbonates of Magnesium.

Second Synthetical Reaction.—To solution of sulphate of magnesium add solution of carbonate of sodium, and boil; the resulting precipitate is *light* carbonate of magnesium (*Magnesiæ Carbonas Levis*, B. P.), a white, partly amorphous, partly minutely crystalline mixture of carbonate and hydrate of

magnesium (3MgCO_3 , Mg_2HO , $4\text{H}_2\text{O}$). A denser, slightly granular precipitate of similar chemical composition (*Magnesia Carbonas*, B. P.) is obtained on mixing strong solutions of the above salts, evaporating to dryness, then removing the sulphate of sodium by digesting the residue in hot water, filtering, washing, and drying the precipitate.



The official proportions for the light carbonate are 10 of sulphate of magnesium and 12 of crystals of carbonate of sodium, each dissolved in 80 of cold water, the solutions mixed, boiled for 15 minutes, the precipitate collected on a filter, well washed, drained, and dried over a water-bath. The heavier carbonate is made with the same proportions of salts, each dissolved in 20 instead of 80 of water, the mixture evaporated quite to dryness, and the residue washed by decantation or filtration until all sulphate of sodium is removed (shown by a white precipitate—sulphate of barium—ceasing to form on the addition of solution of chloride or nitrate of barium to a little of the filtrate). Carbonate of magnesium is of course official in the United States Pharmacopœia (*Magnesii Carbonas*).

Another (Pattinson's) Process.—Considerable quantities of carbonate of magnesium are now prepared by treating dolomite (p. 127) with carbonic acid under pressure. Of the two carbonates the magnesium is dissolved first, and is precipitated from the clear liquid by the heat of a current of steam. (See next reaction.)

Third Synthetical Reaction.—Pass carbonic acid gas, generated as described on page 72, into a mixture of water and carbonate of magnesium contained in a test-tube. After some time, separate undissolved carbonate by filtration; the filtrate contains carbonate of magnesium dissolved by carbonic acid. When of a strength of about 13 grains in one ounce, such a solution constitutes "*Fluid Magnesia*" (*Liquor Magnesie Carbonatis*, B. P.).

Officially 1 pint is directed to be made from freshly prepared carbonate. The latter is obtained by adding a hot

solution of 2 ounces of sulphate of magnesium in half a pint of water to one of $2\frac{1}{2}$ ounces of crystals of carbonate of sodium in another half-pint of water, boiling the mixture for a short time (to complete decomposition), filtering, thoroughly washing the precipitate, placing the latter in 1 pint of distilled water, and transmitting carbonic acid gas through the liquid (say, at the rate of three or four bubbles per second) for an hour or two, then leaving the solution in contact with the gas under slight pressure for twenty-four hours, and, finally, filtering from undissolved carbonate, and, after passing in a little more gas, keeping in a well-corked bottle. Slight pressure is best created by placing the carbonate and water in a bottle fitted with a cork and tubes as for a wash-bottle (p. 102), conveying the gas by the tube which reaches to the bottom, and allowing excess of gas to flow out by the upper tube, the external end of which is continued to the bottom of a common phial containing about an inch of mercury. The phial should be loosely plugged with cotton wool, to prevent loss of metal by spurting during the flow of the gas through it. (Each inch in depth of mercury through which the gas escapes corresponds to about half-a-pound pressure on every square inch of surface within the apparatus.)

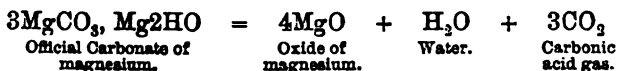
Heat a portion of the solution; true carbonate of magnesium containing combined water ($\text{MgCO}_3, 3\text{H}_2\text{O}$) is precipitated. The water in this compound is probably in the state of water of crystallization, for a salt having the same composition is deposited in crystals by the spontaneous evaporation of the solution of carbonate of magnesium. The official "carbonate" ($3\text{MgCO}_3, \text{Mg}2\text{H}_2\text{O}, 4\text{H}_2\text{O}$) is another of these very common *hydrous* compounds.

Exposed to cold, the solution of "fluid magnesia" sometimes affords large thick crystals ($\text{MgCO}_3, 5\text{H}_2\text{O}$), which, in contact with the air, lose water, become opaque, and then have the composition of those deposited by evaporation ($\text{MgCO}_3, 3\text{H}_2\text{O}$).

Oxide of Magnesium. Magnesia.

Fourth Synthetical Reaction.—Heat light dry carbonate of magnesium in a porcelain crucible over a lamp (or in a larger earthen crucible in a furnace) till it ceases to effervesce on adding, to a small portion, water and acid; the residue is

light magnesia (MgO) (*Magnesia Levis*, B. P.). The same operation on the heavy carbonate yields heavy magnesia (MgO) (*Magnesia*, B. P.). Both are sometimes spoken of as "calcined magnesia." A given weight of the official light magnesia occupies three and a half times the bulk of the same weight of heavy magnesia.



A trace only of magnesia is dissolved by pure water. Moisten a grain or two of magnesia with water, and place the paste on a piece of red litmus-paper; the wet spot, after a time, becomes blue, showing that the magnesia is slightly soluble.

"*Effervescing Citrate of Magnesia*," so-called, is generally a mixture of bicarbonate of sodium, citric acid, tartaric acid, sugar, either carbonate or sulphate of magnesium or both, and flavouring essences. True citrate of magnesium is easily made by heating together calcined magnesia and citric acid; it is frequently prescribed in France in doses of two ounces.

The official Effervescing "Solution of Citrate of Magnesia" (*Liquor Magnesiæ Citratis*, B. P.) is made by dissolving carbonate of magnesium in slight excess of solution of citric acid, adding lemon-syrup, placing the diluted liquid in an aerated-water bottle, dropping in crystals of bicarbonate of potassium, corking, "wiring," and shaking till the crystals are dissolved. This preparation is similar to the *Liquor Magnesiæ Citratis* of the United States Pharmacopœia.

The formula of citrate of magnesium deposited from solution is $\text{Mg}_3\text{C}_6\text{H}_5\text{O}_7, 14\text{H}_2\text{O}$.

REACTIONS HAVING ANALYTICAL INTEREST (TESTS).

First Analytical Reaction.—Add solution of hydrate or carbonate of ammonium to a magnesian solution (sulphate, for example) and warm the mixture in a test-tube; the precipitation of part only of the magnesium as hydrate (Mg_2HO) or carbonate (MgCO_3) occurs. Add now to a small portion of the mixture of precipitate and liquid a considerable excess

of solution of chloride of ammonium; the precipitate is dissolved.

This is an important reaction, especially as regards carbonate of magnesium, the presence of chloride of ammonium enabling the analyst to throw out from a solution barium and calcium by an alkaline carbonate, magnesium being retained. The cause of this retention is found in the tendency of magnesium to form soluble double salts with potassium, sodium, or ammonium. In analysis, the chloride of ammonium should be added before the carbonate, as it is easier to prevent precipitation than to redissolve a precipitate once formed.

Second Analytical Reaction.—To some of the solution resulting from the last reaction, add solution of phosphate of sodium or ammonium; phosphate of magnesium and ammonium (MgNH_4PO_4) is precipitated.—*Third.* To another portion add arseniate of ammonium; arseniate of magnesium and ammonium ($\text{MgNH}_4\text{AsO}_4$) is precipitated.

Note.—Barium and calcium are also precipitated by alkaline phosphates and arseniates. The other precipitants of magnesium are also precipitants of barium and calcium. In other words, there is no *direct* test for magnesium. Hence the analyst always removes any barium or calcium by an alkaline carbonate, as above indicated; the phosphate of sodium, or arseniate or phosphate of ammonium, then become very delicate tests of the presence of magnesium. In speaking of magnesium tests, the absence of barium and calcium salts is to be understood.

QUESTIONS AND EXERCISES.

167. Name the natural sources of the various salts of magnesium.

168. Give a process for the preparation of Epsom salt.

169. Draw diagrams illustrative of the formation of sulphate of magnesium from *magnesite* and from *dolomite*.

170. Show by an equation the process for the preparation of the official Carbonate of Magnesium.

171. What circumstances determine the two different states of aggregation of the official Carbonates of Magnesium (*Magnesia Carbonas* and *Magnesia Carbonas Levis*, B. P.) ?

172. What are the relations of *Magnesia*, B. P., and *Magnesia Levis*, B. P., to the official Carbonates of Magnesium ?

173. How much denser is the one than the other ?

174. Is magnesia soluble in water ?

175. How is "Fluid Magnesia" prepared ?

176. Mention the effects of heat and cold on "Fluid Magnesia."

177. Ascertain how much magnesia (MgO) can be obtained from 100 grains of Epsom salt.

178. Calculate the amount of official Carbonate of Magnesium which will yield 100 grains of magnesia.

179. Can magnesium be detected in presence of barium and calcium ?

180. Describe the analysis of an aqueous liquid containing salts of barium, calcium, and magnesium.

181. How may magnesium be precipitated from solutions containing ammoniacal salts ?

Quantivalence.

On reviewing the foregoing statements regarding compounds of the three univalent radicals, potassium, sodium, and ammonium, and the three bivalent elements, barium, calcium, and magnesium, the doctrine of quantivalence will be more clearly understood and its usefulness be more apparent. Quantivalence, or the value of atoms, is, in short, in chemistry, closely allied to value in commercial barter. A number of articles, differing much in weight, appearance, and general characters, may be of equal money value ; and if these be regarded, for convenience, as having a sort of unit of value, others worth double as much might be termed bivalent, three times as much trivalent, and so on. In like manner, chemical radicals, no matter whether elementary, like potassium (K), iodine (I), or sulphur (S), or compound, like those of nitrates (NO_3), sulphates (SO_4), or acetates ($C_2H_3O_2$), have a given chemical value in relation to each other, and are exchangeable for, or will unite with, each other to an extent determined by that value.

Most chemical salts apparently, though probably not really,

have two parts, a basylous and an acidulous, the one quantitatively balancing the other. The formulæ of the chief of these radicals and their quantivalence are given in the following Table. Examples of formulæ of salts containing univalent, bivalent, and trivalent radicals are also appended.

QUANTIVALENCE OF COMMON RADICALS.

Univalent Radicals, or <i>Monads</i> .		Bivalent Radicals, or <i>Dyads</i> .		Trivalent Radicals, or <i>Triads</i> .	
Acidulous.	Basylous.	Acidulous.	Basylous.	Acidulous.	Basylous.
H	H	O	Ca	PO ₄	As
Cl	K	SO ₄	Mg	BO ₃	Sb
I	Na	CO ₃	Zn	C ₆ H ₅ O ₇	Bi
HO	NH ₄	C ₂ O ₄	Cu	AsO ₃	{ Fe ^{III} (ic) or Fe ^{VI} ₂ (ic)
NO ₃	Ag	C ₄ H ₄ O ₆	Hg(ic)	AsO ₄	
C ₂ H ₃ O ₂	Hg(ous)	S	Fe(ous)	C ₄ H ₃ O ₆	

Note.—Hydrogen (H) as the basylous part of salts has entirely different functions to hydrogen (H) as the acidulous part. Acidulous hydrogen gives compounds commonly termed *hydrides* (e.g., AsH₃); basylous hydrogen is the basylous radical of acids (e.g., HCl, H₂SO₄). On the other hand in compound radicals, e.g., C₂H₃O₂, or NH₄, the properties of hydrogen are no longer apparent; the chemical force resident with the atoms of such radicals seems to be mainly exerted in binding those atoms together.

Examples of Formulæ of Salts containing Univalent, Bivalent, and Trivalent Radicals.

The reader will find instructive practice in writing twenty or thirty imaginary formulæ of salts by placing in juxtaposition acidulous and basylous radicals, as in the following examples. Just as in a pair of scales a 2-lb. weight must be balanced by two 1 lb. weights, or a 4-lb. weight by two 2-lb. weights, or by one 3-lb. and one 1-lb weight, so a bivalent radical unites with a bivalent radical or with two univalent radicals, a quadrivalent radical with two bivalent radicals, or with one trivalent and one univalent radical, and so on.

(R = any basylous radical.) (R = any acidulous radical.)

General formulæ.

Examples.

$R'R'$	. . .	KI, NaCl, $NH_4C_2H_3O_2$, $AgNO_3$.
$R''R'_2$	. . .	$CaCl_2$, $Zn_2C_2H_3O_2$, Pb_2NO_3 ($BaNO_3C_2H_3O_2$).
$R'''R'_3$	. . .	Bi_3NO_3 , AsH_3 , $SbCl_3$.
$R'_2E'_2$	} {	K_2CO_3 , Na_2SO_4 , $H_2C_2H_4O_6$.
$R'R'E''$		$KHCO_3$, $NaHSO_4$, $KN_2C_4H_4O_6$.
R'_3E'''	} {	Am_3PO_4 , $K_3C_6H_5O_7$, H_3AsO_3 .
$R'_2R'E'''$		Na_2HPO_4 , Na_2HASO_4 .
$R''E''$	. . .	$CaCO_3$, MgO , $CuSO_4$, HgO , $FeSO_4$.
$R''_3E'''_3$	. . .	$Ca_3_2PO_4$, $Ca_3_2C_2H_5O_7$.
$R''R'E'''$	. . .	$MgAmPO_4$, $CuHASO_3$.
$R'''E''E$	. . .	$BiONO_3$.
$R'''_2E'''_2E''$	. . .	$Bi_2O_3CO_3$.
$R'''_2E'''_2$	. . .	As_2O_3 , Sb_2O_3 .
$R'''E'''$	. . .	$BiC_2H_5O_7$.
$R'''_2E'_2$	. . .	Fe_2Cl_6 , $Fe_2_6NO_3$, $Fe_2_6C_2H_3O_2$.
$R'''_2E'_2$	. . .	Fe_2O_3 , $Fe_2_3SO_4$.

Quadrivalent Radicals or *Tetrads*, Quinivalent Radicals or *Pentads*, and Sexivalent Radicals or *Hexads*, are known.

EXERCISE.

182. Write an exposition of the doctrine of Quantivalence within the limits of a sheet of note-paper.

DIRECTIONS FOR APPLYING THE FOREGOING ANALYTICAL REACTIONS TO THE ANALYSIS OF AN AQUEOUS SOLUTION OF A SALT OF ONE OF THE METALS, BARIUM, CALCIUM, MAGNESIUM.

Add yellow chromate of potassium to a portion of the solution to be examined; a precipitate indicates barium.

If no barium is present, add chloride and carbonate of ammonium, and boil; a precipitate indicates calcium.

If barium and calcium are proved to be absent, add chloride of ammonium, ammonia, and then either phosphate of sodium or arseniate of ammonium; a white granular precipitate indicates magnesium.

Ammonia is here added to yield the necessary elements to ammonio-magnesian phosphate or ammonio-magnesian arseniate, both of which are highly characteristic precipitates; and chloride of ammonium is added to prevent a mere partial precipitate of the magnesium by the ammonia.

DIRECTIONS FOR APPLYING THE FOREGOING ANALYTICAL REACTIONS
TO THE ANALYSIS OF AN AQUEOUS SOLUTION OF SALTS OF ONE,
TWO, OR ALL THREE OF THE METALS, BARIUM, CALCIUM,
MAGNESIUM.

Add chromate of potassium to the solution; barium, if present, is precipitated. Filter, if necessary, and add to the *filtrate* (that is, the liquid which has run through the filter) chloride, hydrate, and carbonate of ammonium, and boil; calcium, if present, is precipitated. Filter, if requisite, and add phosphate of sodium; magnesium, if present, is precipitated.

Note.—Red chromate of potassium must not be used in these operations, or a portion of the barium will remain in the liquid and be thrown down with, or in the place of, the carbonate of calcium (*vide* p. 111). The yellow chromate must not contain carbonate of potassium, or calcium will be precipitated with, or in the place of, barium. The absence of carbonate is proved by the non-occurrence of effervescence on the addition of hydrochloric acid to a little of the solution of the chromate, previously made hot in a test-tube. If the yellow chromate has been prepared by adding excess of ammonia to solution of red chromate of potassium, its addition to the liquid to be analysed must be preceded by that of solution of chloride of ammonium; the precipitation of a portion of the magnesium (by the free ammonia in the yellow chromate) is thus prevented—for chloride-of-ammonium solution is a good solvent of hydrate (and carbonate) of magnesium, as already stated on page 131.

TABLE OF SHORT DIRECTIONS FOR APPLYING THE FOREGOING ANALYTICAL REACTIONS TO THE ANALYSIS OF AN AQUEOUS SOLUTION OF SALTS CONTAINING ANY OR ALL OF THE METALLIC ELEMENTS HITHERTO CONSIDERED.

To the solution add AmCl , AmHO , Am_2CO_3 ; boil and filter.

Precipitate Ba Ca.		Filtrate Mg Am Na K.	
Wash, dissolve in $\text{HC}_2\text{H}_3\text{O}_2$, add K_2CrO_4 , and filter.		Add Am_2HPO_4 , shake, filter.	
Precipitate Ba.*	Filtrate Ca. Test by $\text{Am}_2\text{C}_2\text{O}_4$.	Precipitate Mg.	Filtrate Am Na K. Evap. to dryness, ignite, dissolve residue in water. Test for K by Pt Cl_4 . Na „ flame. Test. orig. sol. for Am.

Note 1.—The analysis of solutions containing the foregoing metals is commenced by the addition of chloride of ammonium (AmCl) and ammonia (AmHO), simply as a precautionary measure, the former compound preventing partial precipitation of magnesium, the latter neutralizing acids. The carbonate of ammonium (Am_2CO_3), is the important group-reagent—the precipitant of barium and calcium.

Note 2.—In the above, and in subsequent charts of analytical processes, the leading precipitants will be found to be ammonium salts. These, being volatile, can be got rid of towards the end of the operations, and thus the detection of potassium and sodium be in no way prevented—an advantage

* It is perhaps scarcely necessary to state that this precipitate is chromate of barium (BaCrO_4), as any reader who has carefully gone through the “foregoing analytical reactions” will know. The occurrence of chromate of barium at this particular place, however, and under the circumstances described, is abundant evidence of the presence of barium (in some form or other) in the liquid analysed—which was a part of the problem to be solved by the operator. Similar remarks apply of course to the Ca, which is finally precipitated as oxalate (CaC_2O_4), to Mg, which is thrown out as ammonio-phosphate (MgAmPO_4), to Am, Na, and K, and to the elements similarly alluded to in the other subsequent Tables of “short” directions for analysis.

which could not be had if such salts as chromate of potassium or phosphate of sodium were the group-precipitants employed.

Note 3.—Acetic, and not hydrochloric or nitric, acid is used in dissolving the barium and calcium carbonates, because chromate of barium—on the precipitation of which the detection of barium depends—is soluble in the stronger acids, and therefore could not be thrown down in their presence.

Notes on Classification.—The compounds of barium, calcium, and magnesium, like those of the alkali metals, have many analogies; the carbonate, phosphate, and arseniate of each are insoluble in water, which sufficiently distinguishes them from the members of the class first studied. They possess, however, well-marked differences, so that their separation from each other is easy. The solubility of their hydrates in water marks their connection with the alkali metals; the slightness of that solubility, diminishing as we advance farther and farther from the alkalis, baryta being most and magnesia least soluble in water, points to their connection with the next class of metals, the hydrates of which are insoluble in water. These considerations must not, however, be over-valued. Though the solubility of their hydrates places barium nearest and magnesium farthest from the alkali metals, the solubility of their sulphates gives them the opposite order, magnesium-sulphate being most soluble, calcium sulphate next, strontium sulphate third (strontium is a rarer element, which will be mentioned subsequently), and barium-sulphate insoluble in water. These elements are sometimes spoken of as the metals of the alkaline earths.

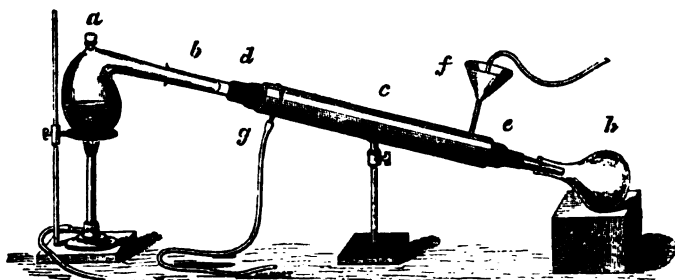
Notes.—In connection with the bivalence of the metals Barium, Calcium, and Magnesium, it is interesting to note that just as bivalent acidulous radicals give salts containing two atoms of univalent basylous radicals, (K_2SO_4 , $NaHSO_4$, H_2CO_3 , $KNaC_4H_4O_6$), so bivalent basylous radicals yield salts containing two atoms of univalent acidulous radicals, as seen in acetonitrates of barium, $BaC_2H_3O_2NO_3$, a salt which is a definite compound, and not a mere mixture of acetate with nitrate of barium. A very large number of such salts is known.

Distillation.

The water with which, in analysis, solution of a salt or dilution of a liquid is effected should be pure. Well- or river-

water are unfit for the purpose, because they contain alkaline and earthy salts (about 20 to 60 grains per gallon), derived from the soil through which the water percolates, and rain-water is not unfrequently contaminated with the dust and *débris* which fall on the roofs whence it is usually collected. Such water is purified by *distillation*, an operation in which the water is by ebullition converted into steam, and the steam condensed again to water in a separate vessel, the fixed earthy and other salts remaining in the vessel in which the water is boiled. On the large scale, ebullition is effected in metal boilers having a hood or head in which is a lateral opening through which passes the steam; on the small scale, either a common glass flask is employed, into the neck of which, by a cork, is inserted a glass tube bent to an acute angle, or a *retort* is used (*a*, fig. 28), a sort of long-necked florence flask,

Fig. 28.



dexterously bent near the body by the glass-worker to an appropriate angle (hence the name *retort*, from *retorqueo*, I bend back). *Condensation* is effected by surrounding the lateral steam-tube with cold water. In large stills the steam-tube, or *condensing-worm*, is usually a metal (tin) pipe, twisted into a spiral form for the sake of compactness, and so fixed in a tub that a few inches of one end of the pipe may pass through and closely fit a hole bored near the bottom of the tub. Cold water is kept in contact with the exterior of the pipe, provision being made for a continuous supply to the bottom, while the lighter water heated by the condensing steam runs off from the top of the column. The condenser for a flask or retort may be a simple glass tube of any size, placed within a much wider tube (a common long, narrow lamp-glass answers very well

for experimental operations), the inner tube being connected at the extremities of the wider by bored corks; a stream of water passes into one end of the enclosed space (the end farthest from the retort), through a small glass tube inserted in the cork, and out at the other end through a similar tube. The common (Liebig's) form of laboratory condenser is a glass tube three-fourths of an inch wide and a yard long (*b*, fig. 28), surrounded by a shorter tin or zinc tube (*c*, fig. 28) two inches in diameter, and having at each extremity a neck, through which the glass tube passes. The ends of the necks of the tin tube, and small portions of the glass tube near them, are connected by means of a strip of sheet caoutchouc carefully bound round, or by short, wide india-rubber tubes (*d* and *e*, fig. 28). An aperture (*f*, fig. 28) near the lower part of the tin tube provides for the admission of a current of cold water, by glass tubing or india-rubber tubing, from the house supply or from a vessel placed above the apparatus; and a similar aperture near the top (*g*, fig. 28) allows the escape of heated water into a vessel or sink. The inner tube may thus constantly be surrounded by cold water, and heated vapours passing through it be perfectly cooled and condensed—and collected in any receiver (*h*, fig. 28).

In distilling several gallons of water for analytical or medicinal purposes (*Aqua Destillata*, B. P.), the first two or three pints should be rejected, because they are likely to contain ammoniacal and other volatile impurities.

Rectification is the process of redistilling a distilled liquid. *Rectified spirit* is a spirit of wine thus treated.

Dry or destructive distillation is distillation in which the condensed products are directly formed by the decomposing influence of the heat applied to the dry or non-volatile substances in the retort or still.

EXERCISE.

183. Write from memory two or three paragraphs descriptive of *distillation*.

RECAPITULATION.

The subject just alluded to (distillation) naturally causes wonder respecting the cause of the physical differences between solid, liquid, and gaseous water. Common observation

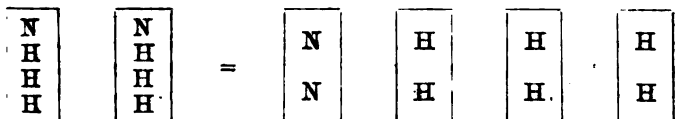
will have suggested to the student that the force of heat has much to do with the differences; and if he will turn to the chapter on *latent heat* in any book on Physics, he will find that as already indicated (p. 88), when ice liquefies by heat a very large amount of heat must be given before the slightest rise of temperature occurs. Afterwards the addition of heat makes the water hotter and hotter until one other point is reached (the boiling-point), when here again a great amount of heat is absorbed without causing the slightest rise in temperature. Afterwards more heat makes the gaseous water hotter and hotter, until, like a bar of iron, the steam, under special conditions, is made red-hot or white-hot. Different bodies absorb different amounts of heat in changing their physical condition from solid to liquid, or liquid to gas (or vapour). The amount is constant for any one body; hence definite comparative numbers may be used for expressing the *latent heats* of substances.

The absorption of heat at particular (liquefying and vaporizing) points must not be confounded with an analogous physical action, namely, the absorption of heat which goes on when a body is rising in temperature. The amount of this absorption, also, differs with different substances. That is to say, if equal weights of several substances, all at the same temperature, be heated to a stated higher temperature, very different amounts of fuel will be required. The particular or specific amount in each case is always the same; hence the *specific heats* of substances may be expressed by numbers. See the chapter on "Specific Heat" in any Manual of Physics.

But after reading what has been stated respecting the constitution of matter (pp. 37 to 41), the chemical student will, in connection with the subject of distillation, be led, once more, to think over the subject of the molecular constitution of solid, liquid, and gaseous water, and of the molecular condition of bodies generally. As previously stated, little can be told him respecting the molecular condition of solids and liquids, for temperature and pressure affects them unequally; whence we conclude, that though the relation to each other of the molecules of any one substance is constant, this relation is different in different bodies. Different *gases*, however, are not differently affected but similarly affected by temperature and pressure; whence we conclude that their molecular constitution—the relation of their molecules to one another—is similar.

Another gas, ammonia, has been brought before the reader since the molecular constitution of gases was considered.

A small quantity of ammonia gas enclosed in the upper part of a roughly graduated test-tube over *mercury* (water would dissolve it) and exposed to the continuous action of the electric spark, by means of wires of platinum fused in the sides of the tube, is decomposed into its elements nitrogen and hydrogen, the bulk of gas operated on being exactly doubled. This expansion is not due to the gaseous molecules receding from each other, but to every two molecules becoming four similar-sized molecules :—



Here each space (rectangular chiefly for convenience in printing) represents a molecule, and each letter an atom. Each space, if regarded as the side of a double cube, may also, for the moment, represent two volumes—such two volumes yielding, in the decomposition, one volume of nitrogen and three volumes of hydrogen, or the four such volumes of ammonia shown in the diagram yielding two volumes of nitrogen and six volumes of hydrogen.

Remembering that a symbol (of a gas) represents one volume, and that a *formula* (of a gas) always represents two volumes, the pupil will now see how full of meaning is such an equation as the following, including, as it does, names of the elements, number of atoms, nature of the molecules, number of the molecules, weights of atoms of the molecules, and therefore weights of bulks of the bodies, or extent of expansion in the disunion of the elements, and therefore their extent of contraction in the act of union :—



AT THIS STAGE THE LEARNER IS AGAIN RECOMMENDED TO READ THE PARAGRAPHS ON THE GENERAL PRINCIPLES OF CHEMICAL PHILOSOPHY (PAGES 29 TO 57), AND TO RETURN TO THEM FROM TIME TO TIME UNTIL THEY ARE THOROUGHLY COMPREHENDED.

ZINC, ALUMINIUM, IRON.

These three elements are classed together for analytical convenience rather than for more general analogies.

ZINC.

Symbol Zn. Atomic weight 65.

Source.—Zinc is tolerably abundant in nature as sulphide (ZnS) or *blende*, and carbonate (ZnCO_3) or *calamine* (from *calamus*, a reed, in allusion to the appearance of the mineral). The ores are roasted to expel sulphur, carbonic acid gas, and some impurities, and the resulting oxide distilled with charcoal, when the metal vaporizes and readily condenses. Zinc is a brittle metal, but at a temperature somewhat below 300°F . is malleable, and may be rolled into thin sheets. Above 400° it is again brittle, and may then be pulverized. At 773°F . it melts, and at a bright red heat is volatile. Zinc in exceptionally fine powder ignites spontaneously, especially if damp, or if stored in a warm place.

Uses.—Its use as a metal is familiar; alloyed with nickel it yields german silver; with twice its weight of copper forms common brass, and as a coating on iron (the so-called *galvanized iron*) greatly retards the formation of rust. Most of the salts of zinc are prepared, directly or indirectly, from the metal (*Zincum*, B. P.).

Quantivalence.—The atom of zinc is bivalent, Zn'' .

Molecular weight.—Some remarks on this point will be made under Mercury.

REACTIONS HAVING (a) SYNTHETICAL AND (b) ANALYTICAL INTEREST.

(a) *Synthetical Reactions.*

Sulphate of Zinc.

First Synthetical Reaction.—Heat zinc (4 parts) with water (20 parts) and sulphuric acid (3 fl. parts) in a test-tube (or larger vessel) until gas ceases to be evolved; solution of sulphate of zinc (ZnSO_4) results. Filter (to separate the particles of lead, carbon, etc., commonly contained in sheet zinc) and concentrate the solution in an evaporating-dish; on

cooling, colourless, transparent, prismatic crystals of Sulphate of Zinc ($\text{ZnSO}_4 \cdot 7\text{H}_2\text{O}$) are deposited (*Zinci Sulphas*, B. P.).



Zinc. Sulphuric acid. Water. Sulphate of zinc. Hydrogen. Water.

Zinc does not displace hydrogen from the sulphuric acid alone, nor from the water alone, yet the mixture affords hydrogen. The probable explanation is that as sulphuric acid combines with several different quantities of water to form definite hydrous compounds (H_2SO_4 , H_2O ; $\text{H}_2\text{SO}_4 \cdot 2\text{H}_2\text{O}$; etc.), it is one of these that is decomposed with elimination of hydrogen. At present we can only say that an unknown (x) amount of water is required in the reaction.

Note.—This reaction affords hydrogen and sulphate of zinc; it also develops electricity. Of several methods of evolving hydrogen, it is the most convenient; of the two or three means of preparing sulphate of zinc, it is that most commonly employed; and of the many reactions which may be utilized in the development of dynamic electricity, it is at present the cheapest and most manageable. The apparatus in which the reaction is effected differs according to the requirements of the operator: if the sulphate of zinc alone is wanted, an open dish is all that is necessary, the action being, perhaps, accelerated by heat; if hydrogen, a closed vessel and delivery-tube; if electricity, square vessels called cells, and certain complementary materials, forming altogether what is termed a battery. In each operation for one product the other two are commonly wasted. It would not be difficult for the operator, as a matter of amusement, to construct an apparatus from which all three products should be collected.

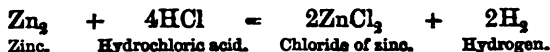
Purification.—Impure sulphate of zinc may be purified in the same manner as impure chloride (see next reaction).

Sulphate of zinc is isomorphous with sulphate of magnesium, and, like that salt, loses six-sevenths of its water of crystallization at 212°F .

Chloride of Zinc.

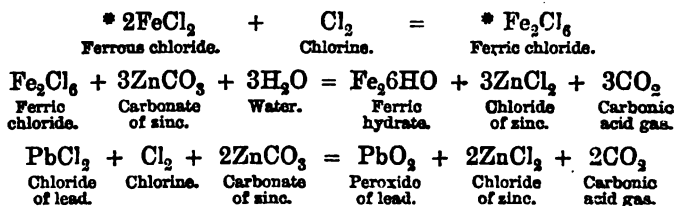
Second Synthetical Reaction.—Dissolve zinc in hydrochloric acid mixed with half its bulk of water; the resulting solution contains chloride of zinc. Evaporate the liquid till no more steam escapes; Chloride of Zinc (ZnCl_2) in a state of fusion remains, and, on cooling, is obtained as an opaque white solid

(*Zinci Chloridum*, B. P.). It is soluble in water, alcohol, or other.

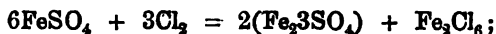


This reaction is analogous to that previously described. The process and product are those of the British Pharmacopœia. Burnett's deodorizing or disinfecting liquid is solution of chloride of zinc.

Purification of Chloride or Sulphate of Zinc.—Zinc sometimes contains traces of iron or lead; and these, like zinc, are dissolved by most acids, with formation of soluble salts: they may be recognised in the liquids by applying the test described hereafter (p. 148) to a little of the solution in a test-tube. Should either be present in the above solution, a little chlorine water is added to the liquid till the odour of chlorine is permanent, and then the whole well shaken with some hydrate of zinc or the common official "carbonate" of zinc (really hydrato-carbonate: see next page). In this way iron is precipitated as ferric hydrate, and lead as peroxide:



In the British Pharmacopœia, the presence of impurities in the zinc is assumed, and the process of purification just described incorporated with the process of preparation of *Zinci Chloridum*, *Liquor Zinci Chloridi*, and *Zinci Sulphas*. In the purification of the sulphate of zinc, the action of chlorine on any ferrous sulphate will result in the formation of ferric sulphate as well as ferric chloride:—



carbonate of zinc will then give chloride as well as sulphate of zinc, and thus the whole quantity of sulphate of zinc be

* It will be noticed that the iron is represented, in these equations, as exerting both bivalent and trivalent activity; this will be alluded to when iron comes under consideration.

slightly contaminated by chloride. On evaporating and crystallizing, however, the chloride of zinc will be retained in the mother-liquor. This process admits of *general* application.

For *Liquor Zinci Chloridi*, B. P., 1 pound of zinc is placed in a mixture of 44 fluid ounces of hydrochloric acid and 20 of water, the mixture ultimately warmed until no more gas escapes, filtered into a bottle, chlorine water added until the liquid after shaking smells fairly of chlorine, about half an ounce or somewhat more of carbonate of zinc shaken up with the solution until a brown precipitate (of ferric hydrate, or peroxide of lead, or both) appears, the whole filtered and the filtrate evaporated to 40 fluid ounces. One fluid ounce contains 366 grains of chloride of zinc. If there is reason to believe that neither iron nor lead is present in the zinc, the treatment with chlorine, water, and carbonate of zinc may be omitted.

Carbonate of Zinc.

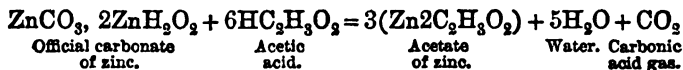
Third Synthetical Reaction.—To solution of any given quantity of sulphate of zinc in twice its weight of water (in a test-tube, evaporating-basin, or other large or small vessel) add about an equal quantity of carbonate of sodium, also dissolved in twice its weight of water, and boil; the resulting white precipitate is so-called Carbonate of Zinc (*Zinci Carbonas*, B. P.), a mixture of carbonate (ZnCO_3) and hydrate (Zn_2HO), in the proportion of one molecule of the former and two of the latter, together with a molecule of water (H_2O). It may be washed, drained, and dried in the usual manner. It is used in the arts under the name of *zinc-white*.



Acetate of Zinc.

Fourth Synthetical Reaction.—Collect on a filter the precipitate obtained in the last reaction, wash with distilled water, and dissolve a portion in strong acetic acid; the resulting solution contains acetate of zinc ($\text{Zn}_2\text{C}_2\text{H}_3\text{O}_2$), and, on evaporating, and setting aside for a day, yields lamellar pearly

crystals ($\text{Zn}_2\text{C}_2\text{H}_3\text{O}_2, 2\text{H}_2\text{O}$). This is the process for *Zinci Acetas*, B. P.



Oxide of Zinc.

Fifth Synthetical Reaction.—Dry the remainder of the precipitated carbonate (by placing the open filter on a plate over a dish of water kept boiling), and then heat it in a small crucible till it ceases to effervesce on the addition of water and acid to trial samples taken out of the crucible from time to time; the product is Oxide of Zinc (*Zinci Oxidum*, B. P.), much used in the form of Ointment (*Unguentum Zinci*, B. P.).

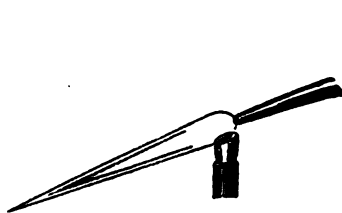


Note.—This oxide is yellow while hot, and of a very pale yellow or slight buff tint when cold, not actually white like the oxide prepared by the combustion of zinc in air. The latter variety occurs in commerce under the name of Hubbuck's oxide of zinc. Its preparation can only be practically accomplished on the large scale, but the chief features of the action may be observed by heating a piece of zinc on charcoal in the blowpipe-flame (fig. 29) till it burns; flocks escape, float about in the air, and slowly fall. These are the old *Flores Zinci*, *Lana Philosophica*, or *Nihilum Album*.

Fig. 29.



Fig. 30.

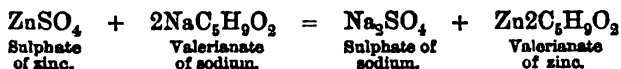


THE BLOWPIPE.

A clear blowpipe-flame consists more or less of two portions (*see* fig. 30), an inner cone at the apex of which are hot gases greedy of oxygen, and an outer cone at the apex of which is excess of hot oxygen. At the latter point oxidizable metals, etc., are readily oxidized, as in the foregoing experiment, and that part of the flame is therefore termed *the oxidizing flame*; in the inner flame oxides and other compounds (a grain of acetate of lead may be employed for illustration) are reduced to the metallic state, hence that part is termed *the reducing flame*. A blowpipe-flame is much altered in character by slight variations in the position of the nozzle of the blowpipe, by the form of the nozzle, by the force with which air is expelled from the blowpipe, and by the character of the jet of gas.

Valerianate of Zinc.

Sixth Synthetical Reaction.—The Valerianate of Zinc ($\text{Zn}2\text{C}_5\text{H}_9\text{O}_2$) (*Zinci Valerianas*, B.P.) is prepared by mixing strong solutions of sulphate of zinc and valerianate of sodium, cooling, separating the white pearly crystalline matter, evaporating at 200°F . to a low bulk, cooling, again separating the lamellar crystals, washing the whole product with a small quantity of cold distilled water, draining and drying by exposure to air at ordinary temperatures. Valerianate of zinc is soluble in ether, alcohol, or hot water.



Note.—The compounds of zinc described in the foregoing six reactions are the only ones mentioned in the British Pharmacopœia; the processes are also those of that work. *Sulphide* and *Hydrate of Zinc* are mentioned in the following analytical paragraphs. The formula of *Sulphite of Zinc* is $\text{ZnSO}_3, 3\text{H}_2\text{O}$.

(b) Reactions having Analytical Interest (Tests).

First Analytical Reaction.—To solution of a zinc salt (sulphate for example), in a test-tube, add solution of sulphhydrate of ammonium (NH_4HS); white sulphide of zinc (ZnS) is

precipitated, insoluble in acetic, but soluble in the stronger acids.

Note.—This is the only white sulphide that will be met with. Its formation, on the addition of the sulphhydrate of ammonium, is therefore highly characteristic of zinc. If the zinc salt contains iron or lead as impurities, the precipitate will have a dark appearance, the sulphides of those metals being black. Hydrate of aluminium, which is also white and precipitated by sulphhydrate of ammonium, is the only substance for which sulphide of zinc is likely to be mistaken, and *vice versâ*; but, as will be seen immediately, there are good means of distinguishing these from each other.

Second Analytical Reaction.—To solution of a zinc salt add solution of ammonia; white hydrate of zinc (Zn_2HO) is precipitated. Add excess of ammonia; the precipitate is redissolved.

This reaction at once distinguishes a zinc salt from an aluminium salt, hydrate of aluminium being, practically, insoluble in ammonia.

Other Analytical Reactions.—The fixed alkali-hydrates afford a similar reaction to that just mentioned, the hydrate of zinc redissolving if the alkali is free from carbonate.——Carbonate of ammonium yields a white precipitate of carbonate and hydrate, soluble in excess.——The fixed alkaline carbonates give a similar precipitate, which is not redissolved if the mixed solution and precipitate be well boiled.——Ferrocyanide of potassium precipitates white ferrocyanide of zinc (Zn_2FeCy_6).

Sulphate of magnesium, which is isomorphous with and indistinguishable in appearance from sulphate of zinc, is not precipitated from its solutions either by ferrocyanide of potassium or sulphhydrate of ammonium.

Antidotes.—There are no efficient chemical means of counteracting the poisonous effects of zinc. Large doses, fortunately, act as powerful emetics. If vomiting has not occurred, or apparently to an insufficient extent, solution of carbonate of sodium (common washing salt), immediately followed by white of egg and demulcents, may be administered.

QUESTIONS AND EXERCISES.

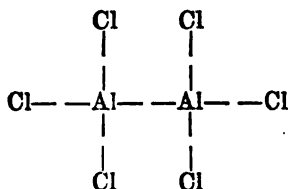
184. Give the sources and uses of metallic zinc.
185. Explain by a diagram what occurs when zinc is dissolved in diluted sulphuric acid.
186. How may solutions of Chloride or Sulphate of Zinc be purified from salts of iron? Give equations descriptive of the reactions.
187. State the formula of the official Carbonate of Zinc, and illustrate by a diagram the reaction which takes place in its production.
188. Give an equation showing the formation of Acetate of Zinc.
189. In what respect does Oxide of Zinc, resulting from the ignition of the carbonate, differ from that produced during the combustion of the metal?
190. How is Valerianate of Zinc prepared?
191. What are the properties of Valerianate of Zinc?
192. Name the more important tests for zinc.
193. How would you distinguish, chemically, between solutions of Sulphate of Zinc and Alum?
194. Describe the treatment in cases of poisoning by salts of zinc.
195. Give reactions distinguishing Sulphate of Zinc from Sulphate of Magnesium.

ALUMINIUM.

Symbol Al. Atomic weight 27.5.

Note.—In the formulæ of aluminium salts, it will be observed that to one atom of metal there are three atoms of other univalent radicals; hence, apparently, the atom of aluminium is trivalent, Al''' . But possibly it is quadrivalent; for one molecule of aluminium compounds includes two atoms of the metal, three-fourths only of whose power may be supposed to be exerted in retaining the other constituents of the molecule, the remaining fourth enabling the aluminium atoms themselves to keep together. This is graphically shown in the following formula of chloride of aluminium (Al_2Cl_6) from Frankland's "Lecture Notes for Chemical Students," which represents each aluminium atom as a body having four arms or bonds, three of which are engaged in grasping the

arms of univalent chlorine atoms, while the fourth grasps the corresponding arm of its brother aluminium atom. Such



graphic formulæ, as they are called, are useful in facilitating the acquirement of hypotheses regarding the constitution of chemical substances, especially if the error be avoided of supposing that they are pictures either of the position or absolute power of atoms in a molecule, or, indeed, the true representation of a molecule at all; for on this point man knows little or nothing.

Source.—Aluminium is very abundant in nature, chiefly as silicate, in clays, slate, marl, granite, basalt, and a large number of minerals. *Mica* or *talc* consists chiefly of silicates of aluminium, iron, and potassium. The sapphire and ruby are almost pure oxide of aluminium. *Rotten stone* is a soft and friable aluminium silicate containing a little organic matter.

The metal aluminium is obtained from the double chloride of aluminium and sodium by the action of metallic sodium, the source of the chloride being the mineral *bauxite*, a more or less ferruginous hydrate of aluminium.

Aluminium-bronze is an alloy of ten parts of aluminium with ninety of copper.

Alum (*Alumen*, B. P., *Aluminii et Potassii Sulphas*, also *Aluminii et Ammonii Sulphas*, U. S. P.), a double sulphate of aluminium and ammonium ($\text{Al}_2\text{SO}_4, \text{Am}_2\text{SO}_4, 24\text{H}_2\text{O}$), may be obtained from aluminous schist (from *σχιστός*, *schistos*, divided), a sort of pyritous slate or shale, by exposure to air; oxidation and chemical change produce sulphate of aluminium, sulphate of iron, and silica, from the silicate of aluminium and bisulphide of iron (iron pyrites) originally present in the shale. The sulphate of aluminium and sulphate of iron are dissolved out of the mass by water, and sulphate or chloride of ammonium added; on concentrating the liquid alum crystallizes out, while the more soluble iron salt remains in the mother-liquor.

Alum is also prepared by directly decomposing the silicate of aluminium in the calcined shale of the coal-measures by hot sulphuric acid, ammonia being added from time to time until a solution strong enough to crystallize is obtained. The liquid well agitated during cooling deposits alum in minute crystals termed *alum-flour*, which is afterwards recrystallized.

Alums.—There are several *alums*, iron or chromium taking the place of aluminium, and potassium or sodium that of ammonium, all crystallizing in an eight-sided form, the octahedron—a sort of double pyramid. They are, apparently, alike in chemical constitution, and their general formula (M = either metal) is M'''_2SO_4 , M'_2SO_4 , $24H_2O$. The alum of the manufacturer commonly occurs in colourless, transparent, octahedral crystals, massed in lumps, which are roughly broken up for trade purposes, but still exhibit the faces of octahedra. It contains sulphate of ammonium or sulphate of potassium, according as one or other is the cheaper.

Preparation of Alum.—Prepare alum by heating a small quantity of powdered pipeclay (silicate of aluminium) with about twice its weight of sulphuric acid for some time, dissolving out the resulting sulphate of aluminium and excess of sulphuric acid by water, and adding ammonia to the clear filtered solution until, after well stirring, the excess of acid is neutralized. (If too much ammonia be added the hydrate of aluminium precipitated when the ammonia is first poured in will not be redissolved on well mixing the whole. Perhaps the readiest indication of neutrality in this and similar cases is the presence of a little precipitate after stirring and warming the mixture.) On evaporating the clear solution crystals of alum are obtained.

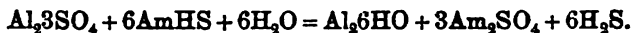
Sulphate of Aluminium or Alum-cake (Al_2SO_4 , $9H_2O$), prepared from natural silicates in the manner just described, is a common article of trade, serving most of the manufacturing purposes for which alum was formerly employed.

Dried alum (*Alumen Exsiccatum*, B.P.) is alum from which the water of crystallization has been expelled by heat, the temperature not exceeding $400^\circ F$. By calculation from the molecular weight of alum, it will be found that the salt contains between 47 and 48 per cent. of water. At temperatures above 400° alum is decomposed, sulphate of ammonium and sulphuric anhydride escaping, and pure oxide of aluminium (Al_2O_3) remaining. Dried alum rapidly re-absorbs water from the atmosphere. It is almost useless as a medicinal preparation.

Roche alum, or *Rock alum* (*roche*, French, rock), is the name of an impure native variety of alum containing iron. The article sold under this name is sometimes an artificial mixture of common alum with oxide of iron.

REACTIONS HAVING ANALYTICAL INTEREST (TESTS).

First Analytical Reaction.—To a solution of an aluminium salt (alum, for example, which contains sulphate of aluminium) add sulphhydrate of ammonium (NH_4HS); a gelatinous white precipitate of hydrate of aluminium falls:—



Second Analytical Reaction.—To solution of alum add ammonia, NH_4HO ; hydrate of aluminium falls: add excess of ammonia; the precipitate is, practically, insoluble.

Principle of Dyeing by help of Mordants.—The precipitated hydrate of aluminium, or alumina, has great affinity for vegetable colouring-matters, and also for the fibre of cloth. Once more perform the above experiment, but before adding the ammonia introduce some decoction of logwood, solution of cochineal, or other similar coloured liquid, into the test-tube. Add now the ammonia, and set the tube aside for the alumina to fall; the latter takes down with it all the colouring principle. In dye works, the fabrics are passed through liquids holding the alumina but weakly in solution, and then through the colouring solutions; from the first bath the fibres abstract alumina, and from the second the alumina abstracts colouring matter. Some other metallic hydrates, notably those of tin and iron, resemble alumina in this property; they are termed *mordants* (from *mordens*, biting); the substances they form with colouring-matters have the name of *lakes*.

Third Analytical Reaction.—To the alum add solution of potash; again hydrate of aluminium falls. Add excess of potash, and agitate; the precipitate dissolves.

Hydrate of aluminium may be precipitated from this solution by neutralizing the potash with hydrochloric acid, and adding ammonia, until, after shaking, the mixture has an

ammoniacal smell, or by adding solution of chloride of ammonium to the potash liquid. But the former way is the better; for it is difficult to know when a sufficiency of the chloride of ammonium has been poured in, whereas reaction with blue and red litmus-paper at once enables the operator to know when excess of hydrochloric acid or of ammonia has been added.

Alkaline carbonates, phosphates, arseniates, and salts of other acidulous radicals also decompose solutions of aluminium salts and produce insoluble compounds of that metal with the several acidulous radicals (except the carbonic), but the resulting precipitates are of no special interest.

QUESTIONS AND EXERCISES.

196. What is there remarkable about the quantivalence of aluminium?

197. Practically what is the quantivalence of the atom of aluminium?

198. Enumerate the chief natural compounds of aluminium?

199. Write down a formula which will represent either of the alums.

200. Which Alum is official, and commonly employed in the arts?

201. State the source and explain the formation of alum.

202. What is the crystalline form of alum? Work a sum showing how much Dried Alum is theoretically producible from 100 pounds of alum. Ans. 52 lb. 6 oz.

203. Show by figures how ordinary ammonium alum is capable of yielding 11.356 per cent. of oxide of aluminium.

204. Why are aluminium compounds used in dyeing?

205. How are salts of aluminium analytically distinguished from those of zinc?

IRON.

Symbol Fe. Atomic weight 56.

Sources.—Compounds of iron are abundant in nature. *Magnetic Iron Ore*, or *Lodestone* (*Lodestons* or *Leadstone*, from the Saxon *ledan*, to lead, in allusion to its use, or, rather, to the use of magnets made from it, in navigation), is the chief ore

from which Swedish iron is made; it is a mixture of ferrous and ferric oxide (FeO , Fe_2O_3). Much of the Russian iron is made from *Specular Iron Ore* (from *speculum*, a mirror, in allusion to the lustrous nature of the crystals of this mineral); this and *Red Hæmatite* (from *αἷμα*, *haima*, blood, so named from the colour of its streak), an ore raised in Lancashire, are composed of ferric oxide only (Fe_2O_3). *Brown Hæmatite*, an oxyhydrate, is the source of much of the French iron. *Spathic Iron Ore* (from *spatha*, a slice, in allusion to the lamellar structure of the ore) is a ferrous carbonate (FeCO_3). An impure ferrous carbonate forms the *Clay Ironstone*, whence most of the English iron is derived. The chief Scotch ore is also an impure carbonate, containing much bituminous matter; it is known as *Black Band*. *Iron Pyrites* (from *πῦρ*, *pur*, fire, in allusion to the production of sparks when sharply struck) (FeS_2) is a yellow lustrous mineral, of use only for its sulphur. As met with in coal it is commonly termed, *coal brasses*. Ferrous carbonate (FeCO_3), chloride (FeCl_2 , $4\text{H}_2\text{O}$), and sulphate (FeSO_4 , $7\text{H}_2\text{O}$) sometimes occur in springs, the water of which is hence termed *chalybeate* (*chalybs*, steel).

Process.—Iron is obtained from its ores by processes of roasting, and reduction of the resulting impure oxide with coal or charcoal in the presence of chalk, the latter uniting with the sand, clay, etc. to form a fusible slag. The *cast iron* thus produced may be converted into *wrought iron* by burning out the 4 or 5 per cent. of carbon, silicon, and other impurities present, by oxidation in a furnace, an operation which is termed *puddling*. *Steel* is iron containing from one to two per cent. of carbon, and is made by the now celebrated *Bessemer* process of burning out from cast iron the variable amount of carbon it contains, and then adding melted iron containing a known proportion of carbon. The official variety of the metal (*Ferrum*, B. P.) "is wrought iron in the form of wire or nails free from oxide": this is the condition in which it is most easily employed for conversion into its compounds. In the form of a fine powder (*see* 17th Reac.) metallic iron is employed as a medicine.

Properties.—The specific gravity of pure iron is 7·844, of the best bar iron 7·7; its colour is bluish white or grey. Bar iron requires the highest heat of a wind-furnace for fusion, but below that temperature assumes a pasty consistence, and in that state two pieces may be joined or *welded* (Germ.

wellen, to join) by the pressure of blows from a hammer. A little sand thrown upon the hot metal facilitates this operation by forming with the superficial oxide of iron a fusible slag, which is dispersed by the blows: the purely metallic surfaces are thus better enabled to come into thorough contact and enter into perfect union. Iron is highly ductile, and of all common metals possesses the greatest amount of tenacity. At a high temperature it burns in the air, forming oxide of iron. *Rust of iron* is chiefly red oxide of iron, with a little ferrous oxide and carbonate; it is produced by action of the moist carbonic acid of the air and subsequent oxidation. Steam passed over scrap iron heated to redness gives hydrogen gas and black oxide of iron. Iron exposed at a high temperature to oxidation by a limited amount of steam (*Barff*) or air (*Bower*), becomes coated with magnetic oxide of so closely coherent and adherent a nature that the metal is permanently protected from alteration by atmospheric and many other influences.

Quantivalence.—Iron combines with other elements and radicals in two proportions; those salts in which the atom of iron appears to possess inferior affinities (in which the other radicals are in the less amount) are termed *ferrous*, the higher being *ferric* salts. In the former the iron exerts bivalent (Fe''), in the latter trivalent activity (Fe''' or Fe_3^{VI}).

The atom of iron is also sometimes considered to be sexivalent, on account of the analogy of its compounds with those of chromium, which is sexivalent, if the formula of its fluoride (CrF_6) be correct, and because the composition of *ferrate of potassium* (K_3FeO_4), a deep-purple salt obtained on passing chlorine through a concentrated solution of potash in which fresh ferric hydrate is suspended, is best explained on the assumption of the sexivalence of its iron.

Why the quantivalence of the atom of iron should vary is not at present known.

The Nomenclature of Iron Salts.—For educational and descriptive purposes the two classes of iron compounds are very conveniently spoken of as *ferrous* and *ferric*, the syllable "*ferr*" common to all indicating their allied ferruginous character, the syllable *ous* and *ic* indicating the lower and higher class respectively—functions fulfilled by these two syllables in other similar cases (sulphurous and sulphuric, mercurous and mercuric). Officially the iron salts are known

by other names, thus, *Sulphate of Iron* (*Ferri Sulphas*) and *Phosphate of Iron* (*Ferri Phosphas*), names which are chemically inexplicit, for there are two sulphates and two phosphates, and the terms do not define which salt is intended. Consistency and uniformity would demand that the names Ferrous Sulphate, Ferrous Phosphate, or similar terms should be employed. Practically, however, the old names cause no confusion, inasmuch as only one sulphate, phosphate, etc., are used in medicine; moreover, the higher salts usually have the prefix *per* attached (as persulphate, perchloride). These names are already well known, can be easily rendered in Latin, and then admit of simple abbreviations and adaptations such as are employed in prescriptions, advantages not possessed by the more rational terms. While, therefore, the comprehension of the chemistry of iron is rendered simple and intelligible by the use of the terms ferrous and ferric, the employment of older and less definite names may very well be continued in pharmacy as being practically more convenient.

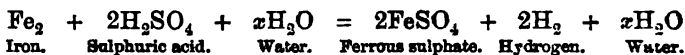
REACTIONS HAVING (a) SYNTHETICAL AND (b) ANALYTICAL INTEREST.

(a) *Synthetical Reactions.*

FERROUS SALTS.

Green Sulphate of Iron. Ferrous Sulphate.

First Synthetical Reaction.—Place iron (small tacks) in sulphuric acid diluted with eight times its bulk of water (in a test-tube, basin, or other vessel of any required size), accelerating the action by heat until effervescence ceases.



The solution contains what is generally known as Sulphate of Iron, that is, Ferrous Sulphate, the lower of the two sulphates, and will yield crystals of that substance ($\text{FeSO}_4 \cdot 7\text{H}_2\text{O}$) (*Ferri Sulphas*, B. P.) on cooling or on further evaporation; or if the hot concentrated solution be poured into alcohol, the mixture being well stirred, the sulphate is at once thrown down in minute crystals (*Ferri Sulphas Granu-*

lata B. P.). At a temperature of 400° F. ferrous sulphate loses six-sevenths of its water, and becomes the *Ferri Sulphas Exsiccata*, B. P. (See the analogous reaction on p. 143.)

Other sources of Ferrous Sulphate.—In the laboratory, ferrous sulphate is often obtained as a by-product in making sulphuretted hydrogen,



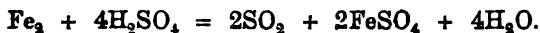
In manufactories it occurs as a by-product in the decomposition of aluminous shale, as already noticed (p. 150).

Ten grains of granulated sulphate of iron dissolved in one ounce of water constitutes "Solution of Sulphate of Iron," B. P. "The solution should be recently prepared."

Notes.—Ferrous sulphate is sometimes termed *green vitriol*. Vitriol (from *vitrum*, glass) was originally the name of any transparent crystalline substance, but afterwards restricted to the sulphates of zinc, iron, and copper, which were, and still are, occasionally known as white, green, and blue vitriol. *Copperas* (probably originally *Copper-rust*, a term applied to verdigris and other green incrustations of copper) is another name for this sulphate of iron, sometimes distinguished as *green copperas*, sulphate of copper being blue copperas. Solid sulphate of iron is a constituent of *Pilula Aloes et Ferri*, B. P. Ferrous sulphate forms a light green double salt with sulphate of ammonium (FeSO_4 , Am_2SO_4 , $6\text{H}_2\text{O}$).

Ferrous sulphate, when exposed to the air, gradually turns brown through absorption of oxygen, ferric oxysulphate ($\text{Fe}_2\text{O}_2\text{SO}_4$) being formed. The latter is not completely dissolved by water, owing to the formation of a still lower insoluble oxysalt ($\text{Fe}_4\text{O}_5\text{SO}_4$) and soluble ferric sulphate $5(\text{Fe}_2\text{O}_2\text{SO}_4) = \text{Fe}_4\text{O}_5\text{SO}_4 + 3(\text{Fe}_2\text{SO}_4)$.

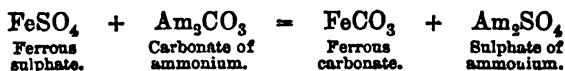
Iron heated with undiluted sulphuric acid gives sulphurous acid gas and ferrous sulphate:—



Carbonate of Iron. Ferrous Carbonate.

Second Synthetical Reaction.—To solution of ferrous sulphate, boiling, in a test-tube, add a solution of carbonate of ammonium (Am_2CO_3) in recently boiled hot water; a white precipitate of ferrous carbonate (FeCO_3) is thrown down, rapidly

becoming light green, bluish green, and, after a long time red, through absorption of oxygen, evolution of carbonic acid gas, and formation of ferric oxyhydrate.



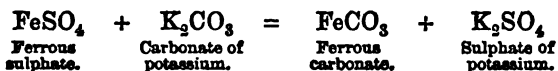
Saccharated Carbonate of Iron.—The above precipitate, rapidly washed with hot well-boiled distilled water, and the moist powder mixed with sugar and quickly dried—in short, all possible precautions taken to avoid exposure to air—forms the saccharated carbonate of Iron (*Ferri Carbonas Saccharata*, B. P.).

The official proportions are two ounces of the sulphate and one ounce and a quarter of the carbonate, each dissolved in half a gallon of hot water; the solutions are mixed and set aside in a deep well-covered pan, the supernatant liquid poured off when the precipitate has subsided, the pan again filled up with boiling water, the liquid once more poured away, the precipitate transferred to a calico filter, drained, gently pressed, and while still somewhat moist rubbed in a mortar with one ounce of sugar, and finally dried over a water-bath.

Carbonate of Iron, mixed with a fourth of its weight of Confection of Roses, forms the *Pilula Ferri Carbonatis*, B. P.

Notes.—The red powder formerly termed Carbonate or Subcarbonate of Iron (*Ferri Carbonas* or *Ferri Subcarbonas*) was ferrous carbonate washed and dried with free exposure to air, the product thus, by the absorption of oxygen and the elements of water and elimination of carbonic acid gas, becoming ferric oxyhydrate, a compound which will come under notice immediately (p. 167). Ferrous carbonate is said to be more easily dissolved in the stomach than any other iron preparation. It is so unstable and prone to oxidation, that it must be washed in water containing no dissolved air and mixed with the sugar (which protects it from oxidation) as quickly as possible. In making the official compound mixture of iron (*Mistura Ferri Composita*, B. P.), "Griffith's mixture," the various ingredients, including the carbonate of potassium, should be placed in a bottle of the required size, space being left for the crystals or solution of ferrous sulphate, which should be added last, the bottle immediately filled up with

the rose-water and securely corked; the minimum of oxidation is thus insured.

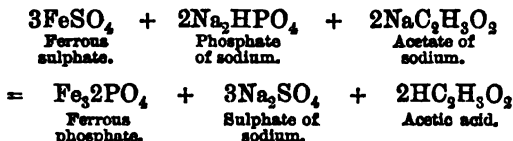


Arseniate of Iron. Ferrous Arseniate.

Third Synthetical Reaction, by which the lower arseniate of iron, ferrous arseniate (*Ferri Arsenias*, B. P.) ($\text{Fe}_3\text{2AsO}_4$), partially oxidized, is formed. This will be noticed again under Arsenicum.

Phosphate of Iron. Ferrous Phosphate.

Fourth Synthetical Reaction.—To solution of ferrous sulphate in a test-tube add a little solution of acetate of sodium, then solution of phosphate of sodium; the lower phosphate of iron, ferrous phosphate ($\text{Fe}_3\text{2PO}_4$) is precipitated (*Ferri Phosphas*, B. P.).



Officially, solutions of 3 ounces of sulphate of iron in a quart of water, and $2\frac{1}{2}$ ounces of phosphate and 1 of acetate of sodium in another quart of water, are well mixed, filtered, the precipitate well washed, and, to prevent oxidation as much as possible, dried at a temperature not exceeding 120° F. These proportions will be found to accord with the molecular weights of the crystalline salts, multiplied as indicated in the foregoing equation. $3(\text{FeSO}_4, 7\text{H}_2\text{O}) = 884$; $2(\text{Na}_2\text{HPO}_4, 12\text{H}_2\text{O}) = 716$; $2(\text{NaC}_2\text{H}_3\text{O}_2, 3\text{H}_2\text{O}) = 272$.

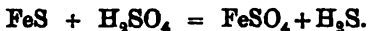
The above reaction also occurs in making *Syrupus Ferri Phosphatis*, B. P. The precipitate should be well washed, or red ferric acetate may be developed after a time.

The use of the acetate of sodium is to insure the absence of free sulphuric acid in the solution—sulphate of sodium being formed together with acetic acid. Sulphuric acid is a powerful solvent of ferrous phosphate; acetic acid is only a weak

solvent. It is impossible to prevent the separation of sulphuric acid, if only ferrous sulphate and phosphate of sodium be employed. The solvent action of the sulphuric acid for the ferrous phosphate is still better prevented by the use of excess of the phosphate of sodium (*Price*), sodium sulphate, and, doubtless, monosodium phosphate (NaH_2PO_4) resulting. Ferrous phosphate is white, but soon oxidizes and becomes slate-blue.

Sulphide of Iron. Ferrous Sulphide.

Fifth Synthetical Reaction.—In a gas-flame or spirit-flame strongly heat sulphur with about twice its weight of iron filings in a test-tube (or in an earthen crucible in a furnace); ferrous sulphide (FeS) is formed. When cold, add water to a small portion, and then a few drops of sulphuric acid; sulphuretted hydrogen gas (H_2S), known by its odour, is evolved.



Sticks of sulphur pressed against a white-hot bar of cast iron give a pure form of ferrous sulphide. The liquid sulphide thus formed is allowed to drop into a vessel of water. (Sulphide of Iron, B. P.). Or melted sulphur may be poured into a crucible full of red-hot iron nails, when a quantity of fluid ferrous sulphide is at once formed and may be poured out on to a slab.

Green Iodide of Iron. Ferrous Iodide.

Sixth Synthetical Reaction.—Place a piece of iodine, about the size of a pea, in a test-tube with a small quantity of water, and add a few iron filings, small nails, or iron wire. On gently warming, or merely shaking if longer time be allowed, the iodine disappears and, on filtering, a clear light-green solution of iodide of iron (FeI_2) is obtained.

The official Ferri Iodidum is formed by gently warming a mixture of 3 parts of iodine, $1\frac{1}{2}$ of fine iron wire, and 12 of distilled water in an iron vessel. When combination is nearly complete (as shown by indications of a sea-green tint) boil for a short time until the whiteness of the froth proves that the iodine has entirely disappeared. The solution is then

filtered and evaporated in a clean bright iron saucepan, ladle, or dish until a drop taken out on the end of an iron wire stirrer solidifies on cooling. The liquid is poured out on a clean smooth slab, broken up and preserved in a glass stoppered bottle. Solid iodide of iron has a crystalline fracture, is "green with a tinge of brown; inodorous, deliquescent, and almost entirely soluble in water, forming a slightly green solution which gradually deposits a coloured sediment and acquires a red colour."

The solid iodide contains about 18 per cent. of water of crystallization and a little oxide of iron. It is deliquescent and liable to absorb oxygen from the air with formation of insoluble ferric oxyiodide or hydrato-iodide. Iodide of iron thus spoilt may be purified by re-solution in water, addition of a little more iodine and some iron, warming, filtering, and evaporating as before.

Ferrous Bromide (FeBr_2), occasionally used in medicine, could be made, as might be expected, in the same way as the iodide.

FERRIC SALTS.

Anhydrous Perchloride of Iron. Ferric Chloride.

Seventh Synthetical Reaction.—Pass chlorine (generated, as usual, from black oxide of manganese and hydrochloric acid in a flask) through sulphuric acid contained in a small bottle

Fig. 31.



PREPARATION OF ANHYDROUS FERRIC CHLORIDE.

and thence by the ordinary narrow glass tubing quite to the bottom of a test-tube containing twenty or thirty small iron tacks (or a florence flask containing 2 or 3 ounces—see fig. 81), the latter kept hot by a gas-flame; the higher chloride of iron, ferric chloride, or the perchloride* of iron (Fe_2Cl_6) is formed and condenses in the upper part of the tube or flask as a mass of small dark iridescent crystals. When a tolerably thick crust of the salt is formed, break off the part of the glass containing it, being careful that the remaining corroded tacks are excluded, and place it in ten or twenty times its weight of water; the resulting solution, poured off from any pieces of glass, is a pure neutral solution of ferric chloride, and will be serviceable in performing analytical reactions.

Precaution.—The above experiment must be conducted in the open air, or in a cupboard having a draught outwards.

Anhydrous Ferrous Chloride.—In breaking up the tube small scales of a light buff colour will be observed adhering to the nails; they are crystals of ferrous chloride (FeCl_2).

Note.—Solution of ferric chloride evolves some hydrochloric acid on boiling, while a darker-coloured solution of ferric oxychloride remains.

Green Chloride of Iron. Hydrous Ferrous Chloride. Solution of Hydrous Ferric Chloride.

Eighth Synthetical Reaction.—Dissolve iron tacks, in a test-tube, in hydrochloric acid; hydrogen escapes, and the solution on cooling, or on evaporation and cooling, deposits *crystallized ferrous chloride* (FeCl_2) associated with four molecules of water of crystallization ($\text{FeCl}_2 \cdot 4\text{H}_2\text{O}$).

Through a portion of the solution of ferrous chloride pass chlorine gas; the ferrous chloride becomes *ferric chloride*.

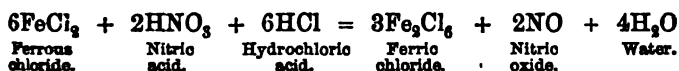
The excess of chlorine dissolved by the liquid in this experiment may be removed by ebullition; but the ferric

* The prefix *per* (and *hyper*) used here and elsewhere is from *ὐπερ*, *hyper*, over or above, and simply means the "highest" of several. Thus perchloride, the highest chloride.

chloride is slightly decomposed at the same time, for the reason just stated. The free chlorine may also be carried off by passing a current of air through the liquid for some time.

Hydrous Ferric Chloride. (*The Official Process.*)

Ninth Synthetical Reaction.—To another portion of the solution of ferrous chloride, in a test-tube, add a little more hydrochloric acid; heat the liquid, and continue to drop in nitric acid until the black colour it first produces disappears; the resulting reddish brown liquid is also solution of ferric chloride.



The black substance is a compound of nitric oxide gas (NO) with a portion of the ferrous salt; it is decomposed by heat.

This is the process for producing the *Liquor Ferri Perchloridi Fortior*, B. P., 2 ounces of iron, 12 fluid ounces of hydrochloric acid, 9 fluid drachms of nitric acid, and 8 ounces of water being employed, and the product boiled down to 10 fluid ounces. Practically it is impossible so to apportion the acids that a solution shall result containing neither excess of acid nor of metal, nor contain ferric nitrate. For most medicinal purposes, however, solution of perchloride of iron containing hydrochloric acid is said to be unobjectionable. On the large scale some time may be saved by adding the acid solution of ferrous chloride to the nitric acid.

Diluted with 3 volumes of water, this strong solution gives the *Liquor Ferri Perchloridi*, B. P.,—or with 3 volumes of rectified spirit the *Tinctura Ferri Perchloridi*, B. P.

Note.—The spirit in the latter preparation is unnecessary, useless, and deleterious; for it acts neither as a special solvent nor as a preservative—the offices usually performed by alcohol (*Tinctura et Succa*, B. P.)—but, unless the liquid contain excess of acid, decomposes the ferric chloride and causes the formation of an insoluble oxychloride of iron. Even if the tincture be acid it slowly loses colour, ferrous chloride and chlorinatted ethereal bodies being formed. The *Liquor*, which

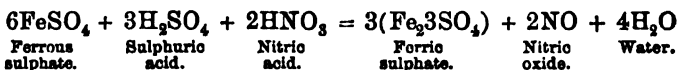
is similar in strength, is doubtless destined to displace the tincture altogether.

Solution of Ferric Chloride evaporated yields a mass of yellow crystals containing $\text{Fe}_2\text{Cl}_6, 12\text{H}_2\text{O}$, or, rarely, red crystals having the formula $\text{Fe}_2\text{Cl}_6, 5\text{H}_2\text{O}$.

The old method of making official solutions of ferric chloride is to dissolve ferric oxide, or hydrate, in hydrochloric acid; but, from the varying character of trade specimens of the ingredients, the liquid is more likely to contain excess or deficiency of iron than the proper proportion.

Persulphate of Iron. Ferric Sulphate.

Tenth Synthetical Reaction.—Dissolve about three-quarters of an ounce of ferrous sulphate and a sixth of its weight of sulphuric acid in an ounce and a half of water in an evaporating-dish, heating the mixture and dropping in nitric acid until the black colour it first produces disappears. The resulting liquid, when made of a certain prescribed strength, is the solution of ferric sulphate, or higher sulphate, "Solution of Persulphate of Iron" of the British Pharmacopoeia, a heavy dark-red liquid.



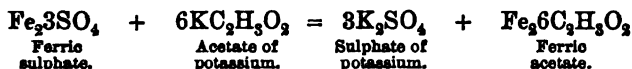
The black colour, as in the previous reaction, is due to a compound of ferrous salt with nitric oxide ($2\text{FeSO}_4 + \text{NO}$).

The definite official solution of ferric sulphate (*Liquor Ferri Persulphatis*, B. P.) is made by adding 6 fluid drachms of sulphuric acid to half a pint of water, warming, dissolving 8 ounces of crystals of sulphate of iron in the liquid, then pouring in nitric acid (6 fluid drachms or rather more) slightly diluted until the mixture turns from a black to a reddish colour, and ruddy nitrous vapours cease to be produced. The whole should measure 11 fluid ounces, being diluted or further evaporated, as the case may be, to this bulk. On the large scale time is saved by adding the mixed fluid to the nitric acid.

Note.—In reactions in which iron passes from ferrous to ferric condition it assumes different properties, the chief being an alteration from bivalent to trivalent activity.

Acetate of Iron. Ferric Acetate.

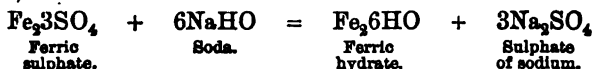
Eleventh Synthetical Reaction.—To a strong solution of ferric sulphate (from which free nitric acid has been removed by evaporating to dryness and redissolving in water) add an *alcoholic* solution of acetate of potassium ($\text{KC}_2\text{H}_3\text{O}_2$), and well shake the mixture; a crystalline precipitate of sulphate of potassium (K_2SO_4) falls, and ferric acetate ($\text{Fe}_26\text{C}_2\text{H}_3\text{O}_2$) remains in solution, forming, when filtered and of definite strength, the *Tinctura Ferri Acetatis*, B. P. The preparation is unstable.



The official proportions are $2\frac{1}{2}$ fluid ounces of "Solution of Persulphate of Iron" with 8 fluid ounces of rectified spirit, mixed with a solution of 2 ounces of acetate of potassium in 10 fluid ounces of spirit, the whole well shaken frequently during an hour, filtered, and the precipitated sulphate of potassium washed by pouring on spirit until the filtrate measures 1 pint. A solution four times this strength, made from ferric hydrate and glacial acetic acid, is stable: it is diluted with spirit as wanted (J. Deane and T. Jeaffreson).

Perhydrate of Iron. Ferric Hydrate.

Twelfth Synthetical Reaction.—Pour a portion of the solution of ferric sulphate into excess of solution of soda; moist ferric hydrate is precipitated (*Ferri Peroxidum Humidum*, B. P.).



Either of the other alkalies (potash or ammonia) will produce a similar reaction; but soda is the one ordered in the British Pharmacopœia, and is cheapest and most convenient.

Ferric hydrate is an antidote to arsenic if administered directly after the poison has been taken.

It converts the soluble arsenic (As_2O_3) into insoluble ferrous arseniate:—



Dried ferric hydrate (then become an oxyhydrate— $\text{Fe}_2\text{O}_3\cdot 2\text{HO}$) (*Ferri Peroxidum Hydratum*, B.P.) has less action on arsenic. Even the moist recently prepared hydrate ($\text{Fe}_2\cdot 6\text{HO}$) loses much of this power as soon as it has become converted into an oxyhydrate ($\text{Fe}_2\text{O}_3\cdot 6\text{HO}$), a change which occurs though the hydrate be kept under water (W. Procter, jun.). According to T. and H. Smith this decomposition occurs gradually, but in an increasing ratio; so that after four months the power of the moist mass is reduced to one-half, and after five months to one-fourth. Now the mere loss of water is not usually followed by any alteration of the essential chemical properties of a compound. It would seem, therefore, that ferric hydrate (two molecules) ($\text{Fe}_2\cdot 12\text{HO}$) probably suffers, on standing, actual decomposition into oxyhydrate ($\text{Fe}_2\text{O}_3\cdot 6\text{HO}$) and water ($3\text{H}_2\text{O}$), and does not merely lose water already existing in it as water. Ferric hydrate is also far more readily soluble in hydrochloric acid, tartaric acid, citric acid, and acid tartrate of potassium, than ferric oxyhydrate. Any formula exhibiting ferric hydrate ($\text{Fe}_2\cdot 6\text{HO}$) as a combination of ferric oxide and water ($\text{Fe}_2\text{O}_3\cdot 3\text{H}_2\text{O}$) is apparently, for these and other reasons, incorrect.

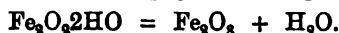
Peroxyhydrate of Iron. Ferric Peroxyhydrate.

Collect the precipitate on a filter, wash, and dry on a plate over hot water; ferric oxyhydrate (*Ferri Peroxidum Hydratum*, B.P.) ($\text{Fe}_2\text{O}_3\cdot 2\text{HO}$) remains. When rubbed to powder it is fit for use in medicine.



Peroxide of Iron. Ferric Oxide.

The above oxyhydrate further decomposes when heated to low redness, ferric oxide (Fe_2O_3) remaining.



The six univalent atoms of the HO, the characteristic elements of all hydrates, are thus, by two successive steps, split up into water and oxygen. But between the hydrate and oxide there obviously may be another oxyhydrate, in which only 2HO is displaced by O''; and such a compound is well known: it is a variety of brown iron ore. The other oxyhy-

drate, $\text{Fe}_2\text{O}_3 \cdot 2\text{H}_2\text{O}$, is also native (needle iron ore), as well as being the *Ferri Peroxidum Hydratum*, B. P.

"Ferri Peroxidum Humidum"	$\text{Fe}'''_2 \text{O}''_4 \cdot 6\text{H}_2\text{O}$
A variety of brown iron ore.	$\text{Fe}'''_2 \text{O}''_4 \cdot 4\text{H}_2\text{O}$
"Ferri Peroxidum Hydratum" (needle ore)	$\text{Fe}'''_2 \text{O}''_3 \cdot 2\text{H}_2\text{O}$
Ferric oxide.	$\text{Fe}'''_2 \text{O}_3$

The moist ferric hydrate, as already stated, when kept for some months, even under water, loses the elements of water, and is converted into an oxyhydrate, having the formula $\text{Fe}_2\text{H}_4\text{O}_9$ (limonite or brown hæmatite), which is either a compound of the above oxyhydrates ($\text{Fe}_2\text{O}_4\text{H}_2\text{O}$) + ($\text{Fe}_2\text{O}_3 \cdot 2\text{H}_2\text{O}$), or is a definite intermediate oxyhydrate ($\text{Fe}_2\text{O}_3 \cdot 6\text{H}_2\text{O}$).

By ebullition with water for seven or eight hours, ferric hydrate is decomposed into water, and an oxyhydrate having the formula $\text{Fe}_2\text{H}_4\text{O}_7$ (Saint-Gilles), which is either a mixture of the official oxyhydrate ($\text{Fe}_2\text{O}_3 \cdot 2\text{H}_2\text{O}$) with ferric oxide (Fe_2O_3), or a definite intermediate body ($\text{Fe}_2\text{O}_3 \cdot 2\text{H}_2\text{O}$). The relation of these bodies to each other will be apparent from the following Table, in which, for convenience, the formulæ of ferric hydrate and oxide are doubled:—

Ferric hydrate (B. P.) (as stalactite)	$\text{Fe}_2 \text{O}_3 \cdot 12\text{H}_2\text{O}$
Kilbride mineral (?)	$\text{Fe}_2 \text{O}_3 \cdot 10\text{H}_2\text{O}$
Brown iron ore (Hüttenrode and Raschau)	$\text{Fe}_2 \text{O}_3 \cdot 8\text{H}_2\text{O}$
Old, or frozen, ferric hydrate (limonite).	$\text{Fe}_2 \text{O}_3 \cdot 6\text{H}_2\text{O}$
Ferric oxyhydrate (B. P.) (gothite)	$\text{Fe}_2 \text{O}_3 \cdot 4\text{H}_2\text{O}$
Boiled ferric hydrate (turgite).	$\text{Fe}_2 \text{O}_3 \cdot 2\text{H}_2\text{O}$
Ferric oxide (red hæmatite)	$\text{Fe}_2 \text{O}_3$

A ferric oxycarbhydrate ($\text{Fe}_2\text{OCO}_3 \cdot 8\text{H}_2\text{O}$) has been obtained by Rother.

The official ferric oxyhydrate ($\text{Fe}_2\text{O}_3 \cdot 2\text{H}_2\text{O}$), termed in the Pharmacopœia *Hydrated Peroxide of Iron*, under the assumption that it is a compound of ferric oxide and water ($\text{Fe}_2\text{O}_3 \cdot \text{H}_2\text{O}$), was formerly made by mixing solutions of ferrous sulphate and carbonate of sodium and exposing the resulting ferrous carbonate to the air until it was nearly all converted into ferric oxyhydrate; hence its old name, still sometimes seen on old bottles, of *Ferri Carbonas* and *Ferri Subcarbonas*.

Ferric Oxide. (Another Process.)

Thirteenth Synthetical Reaction.—Roast a crystal or two of

ferrous sulphate in a small crucible until fumes cease to be evolved; the residue is a variety of ferric oxide (Fe_2O_3), or peroxide of iron, known in trade as *red oxide of iron*, *colcothar*, *crocus*, *rouge* (mineral), *Venetian red*, etc. It has sometimes been used in pharmacy in mistake for the official oxyhydrates (*vide* 12th Synthet. Reac.), from which it differs not only in composition but in the important respect of being almost insoluble in acids.

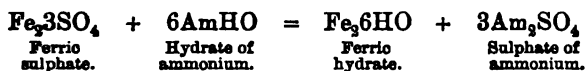
The "Scale" Compounds of Iron.

Fourteenth Synthetical Reaction.—Repeat the twelfth reaction, introducing a little solution of citric or tartaric acid or acid tartrate of potassium, before adding to the alkali (soda, potash, or ammonia), and notice that now no precipitation of ferric hydrate occurs. This experiment serves to illustrate not the manufacture of a scale compound but the chemistry of the manufacture. The effect is due to the formation of double compounds, termed Ammonio-Citrate, Potassio-Citrate, Ammonio-Tartrate, Potassio-Tartrate, and similar sodium compounds of iron, which remain in solution along with the secondary product—sulphate of the alkali metal. Such ferric compounds, made with certain prescribed proportions of recently prepared ferric hydrate (from which all alkaline sulphate has been washed), and the respective acids (tartaric or citric) or acid salts (acid tartrate of potassium), etc., and the solutions evaporated to a syrupy consistence and spread on flat plates till dry, form the scale preparations known as *Ferri et Ammoniae Citras*, B. P., and *Ferri Potassio-tartras*, or, rather, *Ferrum Tartaratum*, B. P. A mixture of ferric citrate with citrate of ammonium and citrate of quinine yields, by similar treatment, the well-known scales of *Ferri et Quiniae Citras*, B. P.

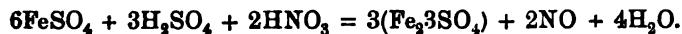
Specimens of these substances may be prepared by attending to the following details. It is essential, first, that the ferric hydrate be thoroughly washed, or an insoluble oxysulphate will be formed; second, that the ferric hydrate be rapidly washed, or an insoluble ferric oxyhydrate will be

produced; thirdly, that the whole operation be conducted quickly, or reduction to green ferrous salt will occur; fourthly, that the solutions of the salts be not evaporated at a higher temperature than that stated, or decomposition will take place; and, fifthly, that excess of ferric hydrate be employed.

In the pharmacopœial processes for the three scale compounds, the ferric hydrate is in each case freshly made from solution of ferric sulphate by precipitation with solution of ammonia:—



the solution of ferric sulphate being made of a definite strength from a known weight of ferrous sulphate. The reason for adopting this course is, that ferric hydrate is unstable and cannot be weighed, because it cannot be dried without decomposing and becoming insoluble, as explained under the 12th reaction. This definite solution of ferric sulphate (*Liquor Ferri Persulphatis*, B. P.) is made by adding six fluid drachms of sulphuric acid to half a pint of water, warming, dissolving eight ounces of crystals of sulphate of iron in the liquid, pouring in nitric acid (six fluid drachms or rather more) slightly diluted until the mixture turns from a black to a reddish colour, and ruddy nitrous vapours cease to be produced; the whole should measure eleven fluid ounces, being diluted or further evaporated, as the case may be, to this bulk.



Ferri et Ammonia Citras, B. P. (*Ferri et Ammonii Citras*, U. S. P.).—Ferric hydrate is dissolved in solution of citric acid, ammonia added, and the whole evaporated to dryness.

To prepare the ferric hydrate, dilute eight (or, rather, nine and a half) fluid ounces of the above solution of ferric sulphate with about a quart of water; pour this into two or three pints of water containing excess of solution of ammonia—about five fluid ounces of “Strong Solution of Ammonia,” or fifteen ounces of “Solution of Ammonia.” (If the opposite course were adopted, the alkaline liquid poured into the ferric solution, the precipitate would contain ferric oxysulphate, or hydrato-sulphate, which interferes with the brilliancy

of the scales.) Thoroughly stir the mixture (it will smell strongly of ammonia if enough of the latter has been added), allow the precipitate to subside, pour away the supernatant liquid, add more water, and repeat the washing until a little of the liquid tested for by-product (sulphate of ammonium) by solution of chloride or nitrate of barium ceases to give a white precipitate (sulphate of barium). Collect the ferric hydrate on a filter, drain, and place, while still moist, in a solution of four ounces of citric acid in eight of water, placed in an evaporating-basin over a water-bath; stir frequently, until nearly the whole of the hydrate has dissolved. To the solution, when cold, add nearly two fluid ounces of strongest (or five and a half of weak) solution of ammonia, filter, evaporate over a water-bath to the consistence of syrup, thinly spread on panes of glass, and dry (at a temperature not exceeding 100°F.). The product scales off the glass in deep red transparent laminæ.

Ferri et Quiniæ Citras, B.P.—Ferric hydrate and pure quinia are dissolved in solution of citric acid, ammonia added, and the whole evaporated to dryness. The product contains citrate of iron, citrate of quinia, and citrate of ammonium.

The ferric hydrate is obtained from four and a half fluid ounces of the solution of ferric sulphate, with all the precautions described in the previous paragraph, a proportionate quantity of ammonia being employed.

While the ferric hydrate is being washed, prepare the quinia by dissolving one ounce of the ordinary sulphate of quinia in eight ounces of distilled water, acidified with sufficient sulphuric acid to dissolve the sulphate (about twelve fluid drachms of the official "diluted sulphuric acid"), and to the clear liquid add solution of ammonia, well mixing the product by stirring, until the whole of the quinia is precipitated (that is, until the mixture, after thorough agitation, smells of ammonia). Collect the precipitate on a filter, let it drain, and wash away adhering solution of sulphate of ammonium by passing through it about a pint and a half of distilled water.

(It will be observed that the principle involved in the preparation of quinia from its sulphate is identical with that which obtains in the precipitation of alumina, ferric hydrate, or hydrate of zinc, etc. Such a soluble sulphate—or, indeed, any similar soluble salt—has its acidulous constituent removed by the superior affinity of the basylous radical in ammonia, or other alkali, an insoluble precipitate and a new soluble sulphate being formed. The latter is washed away, leaving the former

pure. In such manipulations, when economy has to be practised, soda is the alkali generally employed. Ammonia, however, has the advantage of showing the moment when its work of removing an acidulous radical is completed; for the salts which ammonium forms with such acidulous radicals as SO_4 , Cl , NO_3 , and $\text{C}_2\text{H}_3\text{O}_2$ are inodorous, while ammonia has a powerful odour; so long, therefore, as the salt to be decomposed is not wholly attacked, the addition of ammonia does not give an ammoniacal odour to the mixture, the ammonia, as such, being, in fact, destroyed; but when the work is accomplished, the quantity of ammonia last added remains as ammonia, and communicates its natural smell to the liquid.)

The ferric hydrate and quinia being now washed and drained, dissolve the former, and afterwards the latter, in a solution of three ounces of citric acid in five of distilled water, the acid liquid being warmed over a water-bath, and portions of the precipitates stirred in as fast as solution is effected. "Let the solution cool, then add, in small quantities at a time, twelve fluid drachms of solution of ammonia diluted with two fluid ounces of distilled water, stirring the solution briskly, and allowing the quinia which separates with each addition of ammonia to dissolve (in the acid) before the next addition is made (excess of ammonia must be avoided or the quinia will be precipitated). Filter the solution, evaporate to the consistence of a thin syrup, and then dry in thin layers on flat porcelain or glass plates at a temperature of 100° . Remove the dry salt in flakes, and keep it in a stoppered bottle." Long-continued exposure to sunlight causes opacity in the scales, and renders them difficultly soluble (Wood).

Ferrum Tartaratum, B.P. (*Ferri et Potassii Tartras*, U.S.P). — Ferric hydrate is dissolved in solution of acid tartrate of potassium, and the whole evaporated to dryness.

The ferric hydrate obtainable from five and a half fluid ounces (six is better) of the official solution of ferric sulphate by the action of ammonia, in the manner detailed in the previous paragraphs, is mixed (in a mortar), while still moist but well drained, with two ounces of acid tartrate of potassium. The whole is set aside for about twenty-four hours, occasionally triturated to promote contact and reaction of the molecules (otherwise somewhat sluggish in attacking each other), and then heated in a dish over a water-bath to a temperature not exceeding 140°F .; a pint of distilled water is then added, and the mixture kept warm until nothing more will dissolve; filtered, evaporated at a temperature not exceeding 140° (greater heat causes decomposition), and, when the mixture has the consistence of syrup, spread on panes of glass

and allowed to dry (in any warm and light place shown by a thermometer to be not hotter than 140°). The dry salt is thus obtained in flakes. It should be kept in well-closed bottles.

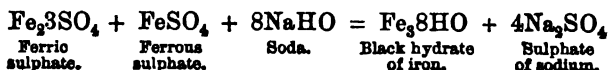
The foregoing are the only official scale preparations of iron. Many others of similar character might be formed. The *Citrate* dissolves slowly in cold but readily in warm water. None crystallize or give other indications of definite chemical composition. Their properties are only constant so long as they are made with unvarying proportions of constituents. Want of chemical compactness, the loose state in which the iron is combined, precludes their recognition as well-defined chemical compounds, yet possibly enables them to be more readily assimilated as medicines than some of the more definite ferrous and ferric salts. A definite *ferrous tartrate* ($\text{FeC}_4\text{H}_4\text{O}_6$) and *ferrous citrate* ($\text{FeHC}_6\text{H}_5\text{O}_7$, H_2O) have been obtained by reaction of iron and acid in hot water. They occur as white gritty masses of microscopic crystals. In the United States Pharmacopœia *Ferri et Ammonii Tartras* is included.

Wine of Iron, or "Steel" wine (*Vinum Ferri*, B. P.), made by digesting iron wire in sherry wine, probably contains tartrate of potassium and iron and other iron salts, formed by action of the metal on the acid tartrate of potassium and tartaric, citric, malic, and acetic acids present in the wine. *Vinum Ferri Oitratis*, B. P., is a solution of ammonio-citrate of iron in orange wine.

Black Hydrate of Iron. Ferroso-ferric Hydrate.

Ferri Oxidum Magneticum, B. P.

Fifteenth Synthetical Reaction.—To two-thirds of a small quantity of a solution of ferrous sulphate add a little sulphuric acid; warm, and gradually add nitric acid, as described in the tenth reaction, care being taken not to allow one drop more nitric acid than necessary to fall into the test-tube. Add the other third of ferrous sulphate, shake, and pour the liquid into excess of an alkali; black (at first brown) hydrate of iron, or ferroso-ferric hydrate ($\text{Fe}_38\text{HO} = \text{Fe}_2\text{HO}, \text{Fe}_6\text{HO}$),



is produced. It is so readily attracted by a magnet, even when moist, as to collect round the poles when the instrument is immersed in the supernatant liquid. Hence the B. P. name, *Ferri Oxidum Magneticum*.

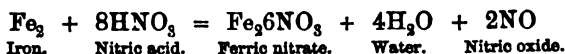
In this process the nitric acid oxidizes the hydrogen of the sulphuric acid, the sulphuric radical uniting with the ferrous sulphate, the iron of which is at the same time altered from the ferrous to the ferric condition, ferric sulphate being formed. If too much nitric acid be employed, the second portion of ferrous sulphate will also be converted into ferric salt, and the solution, on the addition of alkali, yield only red ferric hydrate. This result may be avoided by evaporating the solution of ferric sulphate nearly to dryness, thus boiling off excess of nitric acid, or by pouring first the ferric and then the ferrous liquid into the alkali and thoroughly stirring the mixture; any nitric acid is then neutralized and rendered incapable of oxidizing the ferrous sulphate subsequently added.

Black hydrate of iron is decomposed by heat, yielding, in a closed vessel, oxyhydrates and, finally, black oxide of iron or ferroso-ferric oxide. Heated in the air it absorbs oxygen and gives ferric oxide. The black *forge-scales* which collect near the blacksmith's anvil have the composition of ferroso-ferric oxide; the black magma formed on exposing a mixture of iron and water to the air is ferroso-ferric hydrate; but these varieties are apt to contain particles of metal and, hence, give hydrogen gas when dissolved in acids—a character which distinguishes them from the official preparation.

If a dried specimen of the black hydrate of iron be required, the mixture should be well boiled and then set aside for an hour or two to favour aggregation of the particles, the mixture filtered, and the precipitate washed until the washings contain no trace of sulphate (that is until they no longer yield a white precipitate with chloride of barium). Black hydrate of iron absorbs oxygen even at the temperature of the water-bath; it should consequently be dried at 120° , a temperature at which only slight oxidation occurs.

Pernitrate of Iron. Ferric Nitrate.

Sixteenth Synthetical Reaction.—Place a few iron tacks in dilute nitric acid and set aside; solution of ferric nitrate, or pernitrate of iron, is formed ($\text{Fe}_2\text{6NO}_3$).

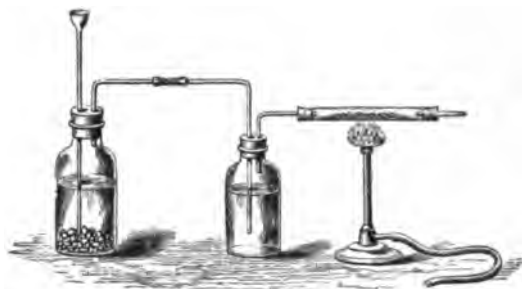


This solution, made with care and of a prescribed strength, forms the *Liquor Ferri Pernitratis*, B. P. The process is as follows:—Four and a half fluid ounces of nitric acid are diluted with sixteen ounces of distilled water, and one ounce of iron wire, free from rust, dissolved in the mixture, the latter being kept cool to avoid violence of action. The liquid is finally filtered and diluted to thirty fluid ounces.

Ferric nitrate and ferric acetate unite to form various acetonitrates, amongst which is one having the formula $\text{Fe}_24\text{C}_2\text{H}_3\text{O}_2\text{NO}_3\text{HO}$, $4\text{H}_2\text{O}$ crystallizing in hard, shining, brownish-red prisms.

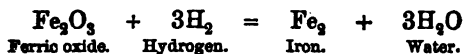
Reduced Iron.

Fig. 32.



PREPARATION OF REDUCED IRON.

Seventeenth Synthetical Reaction.—Pass hydrogen gas (dried by passing over pieces of chloride of calcium contained in a tube, or through sulphuric acid in a wash-bottle) over a small quantity of ferric oxyhydrate or oxide contained in a tube arranged horizontally (a test-tube the bottom of which has been accidentally broken serves very well), the oxide being kept hot by a gas-flame; oxygen is removed from the oxide by the hydrogen, steam escapes at the open end of the tube, and after a short time, when moisture ceases to be evolved, metallic iron, in a minute state of division, remains. (See fig. 32.)



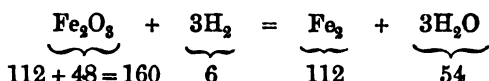
While still hot throw the iron out into the air; it takes fire and falls to the ground as oxide.

If the ferric oxide is reduced in a gun-barrel heated by a strong furnace, the particles of iron aggregate to some extent, and, when cold, are only slowly oxidized in dry air. This latter form of reduced iron is *Fer réduit*, or *Quevenne's Iron*, the *Ferri pulvis*, or *Ferrum Redactum*, B. P.—“a fine greyish-black powder, strongly attracted by the magnet, and exhibiting metallic streaks when rubbed with firm pressure in a mortar.” It is often administered in the form of lozenges (*Trochisci Ferri Redacti*, B. P.), gum and sugar protecting the iron from oxidation as well as forming a vehicle for its administration.

Note I.—The spontaneous ignition of the iron in the above experiment is an illustration of the influence of minute division on chemical affinity. The action is the same as occurs whenever iron rusts, and the heat evolved and amount of oxide formed is not greater from a given quantity of iron; but the surface exposed to the action of the oxygen of the air is, in the case of this variety of reduced iron, so enormous compared with the weight of the iron, that heat cannot be conducted away sufficiently fast to prevent elevation of temperature to a point at which the whole becomes incandescent. In the slow rusting of iron, escape of heat occurs, but is not observed, because spread over a length of time; in the spontaneous ignition of reduced iron the whole is evolved at one moment. The mixture of lead and carbon (lead pyrophorus), resulting when tartrate of lead is heated in a test-tube until fumes cease to be evolved, spontaneously ignites when thrown into the air, and for the same reason. Many substances, solid and liquid, if sufficiently finely divided and liable to oxidation, and especially if exposed in a warm place, become hot and even occasionally spontaneously burst into flame. Oil on cotton waste, powdered charcoal, coal, especially if pyritic or if very porous or if powdered, resins in powder, and even flour, are familiar illustrations of materials liable to “heat” or even burn spontaneously.

Note II.—The student having time and opportunity for the experiment is advised to make this seventeenth reaction a roughly quantitative one, by way of realizing what has been

stated (see, again, the General Principles of Chemical Philosophy, pp. 29–55) respecting the action of chemical force on definite weights only of matter. Three tubes, similar to the oxide-tube shown in the engraving, should be prepared, the second being connected to the first and the third to the second by india-rubber tubing in the usual manner. The first tube should contain pieces of chloride of calcium to absorb any traces of moisture not retained by the sulphuric acid. The second tube (the ends of the small tube being temporarily closed by small corks) should be weighed in any ordinary scales which will turn with a quarter of or half a grain, and, the weight being noted, 160 grains of dry ferric oxide should be neatly placed in the middle of the tube. (The oxide must be previously gently heated in a small crucible over a lamp to remove all traces of moisture.) The third tube should contain pieces of chloride of calcium to absorb the water produced in the reaction, and just before being connected should be weighed. The operation is now carried out. At its close, and when the middle tube is cold, the latter tube and the third tube are again weighed. The oxide-tube should weigh 48 grains less than before, and the terminal tube 54 grains more than before.



The operation is more quickly and easily performed if one-half or one-quarter of the weight of oxide be taken; in that case one-half or one-quarter of the weights of iron and of water will be obtained. Indeed any weight of oxide may be employed; the amount of iron and water resulting will be *always exactly proportionate* to the weights just mentioned. Thus 16 parts of oxide yield 11·2 of iron and 5·4 of water. Iron, hydrogen, and oxygen always combine in proportions of 56, 1, and 16 respectively. Such facts justify us in agreeing that the symbol Fe shall stand for 56 parts by weight of iron, H for 1 part by weight of hydrogen, and O for 16 parts by weight of oxygen.

(b) *Reactions having Analytical Interest (Tests).*

(The iron occurring as a ferrous salt.)

First Analytical Reaction.—Pass sulphuretted hydrogen

(H_2S) through a solution of a ferrous salt (e.g. ferrous sulphate) slightly acidulated by hydrochloric acid; no precipitate occurs.

This is a valuable negative fact, as will be evident presently.

Second Analytical Reaction.—Add sulphhydrate of ammonium (NH_4HS) to solution of a ferrous salt; a black precipitate of ferrous sulphide (FeS) falls.



Third Analytical Reaction.—Add solution of ferrocyanide of potassium (yellow prussiate of potash), $K_4Fe^{''}Cy_6$, or $K_4Fcy^{''}$, to solution of a ferrous salt; a precipitate ($K_2Fe^{''}Fe^{''}Cy_6$, or $K_2Fe^{''}Fcy$) falls, at first white or bluish grey, but rapidly becoming blue, owing to absorption of oxygen.

Fourth Analytical Reaction.—To solution of a ferrous salt add ferridcyanide of potassium (red prussiate of potash, $K_6Fe^{'''}Cy_{12}$, or K_6Fdcy ; a precipitate ($Fe^{'''}_3Fe^{''}_2Cy_{12}$, or $Fe^{'''}_3Fdcy$) resembling prussian blue (Turnbull's Blue) is thrown down.

Other Analytical Reactions.—The precipitates produced from ferrous solutions on the addition of alkaline carbonates, phosphates, and arseniates, as already described in the synthetical reactions of ferrous salts, are characteristic, and hence have a certain amount of analytical interest, but are inferior in this respect to the four reactions above mentioned.

Note.—Alkalies (potash, soda, or ammonia) are incomplete precipitants of ferrous salts, hence are almost useless as tests. To solution of a ferrous salt add ammonia (NH_4HO); on filtering off the whitish ferrous hydrate and testing the solution with sulphhydrate of ammonium, iron will still be found. To another portion of the ferrous solution add a few drops of nitric acid and boil; this converts the ferrous into ferric salt, and now alkalies will wholly remove the iron, as already twice seen during the performance of the synthetical experiments.

In actual analysis, the separation of iron as ferric hydrate is an operation of frequent performance. This is always

accomplished by the addition of alkali, and, if the iron occurs as a ferrous salt, by previous ebullition with a little nitric acid. Ferrocyanide and ferridcyanide of potassium are the reagents used in distinguishing ferrous from ferric salts.

(The iron occurring as a ferric salt.)

Sixth Analytical Reaction.—Through a ferric solution (ferric chloride *e.g.*) pass sulphuretted hydrogen; a white precipitate of the sulphur of the sulphuretted hydrogen falls, and the ferric is reduced to a ferrous salt, the latter remaining in solution. This reaction is of frequent occurrence in practical analysis.



Seventh Analytical Reaction.—Add sulphhydrate of ammonium to a ferric solution; the latter is reduced to the ferrous state, and black ferrous sulphide (FeS) is precipitated as in the second analytical reaction, sulphur being set free.

Eighth Analytical Reaction.—To a ferric solution add ferrocyanide of potassium (K_4FeCy_6 , or $\text{K}_4\text{Fcy}'''$); a precipitate of prussian blue (the common pigment) occurs ($\text{Fe}'''\text{Fe}''\text{Cy}_6$, or $\text{Fe}'''\text{Fcy}'''_2$).

Ninth Analytical Reaction.—To a ferric solution add solution of ferridcyanide of potassium; no precipitate occurs, but the liquid is darkened to a greenish or olive hue, according to the strength.

Tenth Analytical Reaction.—This is the production of a red precipitate of ferric hydrate, on the addition of alkalies to ferric salts, and is identical with the twelfth synthetical reaction.

Note.—This reaction illustrates the conventional character of the terms synthesis and analysis. It is of equal importance to the manufacturer and the analyst, and is synthetical or analytical according to the intention with which it is performed.

Other ferric reactions have occasional analytical interest. In neutral ferric solutions the tannic acid in aqueous infusion of galls occasions a bluish-black inky precipitate, the basis of

most black writing-inks.—(The *Mistura Ferri Aromatica* of the British Pharmacopœia, made by digesting metallic iron in an infusion of various vegetable substances, contains taunate, or rather tannates of iron; it is commonly known in Ireland by the name of Heberden's Ink, after the physician by whom it was first used. It contains about 1 grain of iron in 1 pint.)

—*Sulphocyanate of Potassium* (KCyS) causes the formation of ferric sulphocyanate, which is of a deep blood-red colour.

—There is no ferric carbonate; alkaline carbonates cause the precipitation of ferric hydrate, while carbonic acid gas escapes.

Note.—Cyanogen (CN, or Cy'), ferro-cyanogen (FeC_6N_6 , or FeCy_6 , or simply Fcy'''), and ferridcyanogen ($\text{Fe}_3\text{Cy}_{12}$, or Fdcy'') are radicals which play the part of non-metallic elements, just as ammonium in its chemical relations resembles the metallic elements. They will be again referred to.

Memorandum.—The reader must on no account omit to write out equations or diagrams expressive of each of the reactions of iron, analytical as well as synthetical. It is presumed that this has already been done immediately after each reaction has been performed.

DIRECTIONS FOR APPLYING THE FOREGOING ANALYTICAL REACTIONS
TO THE ANALYSIS OF AN AQUEOUS SOLUTION OF SALTS CONTAINING ONE OF THE METALS, ZINC, ALUMINIUM, IRON.

Add solution of ammonia gradually :—

A dirty-green precipitate indicates iron in the state of a ferrous salt.

A red precipitate indicates iron in the state of a ferric salt.

A white precipitate, insoluble in excess, indicates the presence of an aluminium salt.

A white precipitate, soluble in excess, indicates zinc.

These results may be confirmed by the application of some of the other tests to fresh portions of the solution.

TABLE OF SHORT DIRECTIONS FOR APPLYING THE FOREGOING ANALYTICAL REACTIONS TO THE ANALYSIS OF AN AQUEOUS SOLUTION OF SALTS OF ONE, TWO, OR ALL THREE OF THE METALS, ZINC, ALUMINIUM, IRON.

Boil about half a test-tubeful of the solution with a few drops of nitric acid. This insures the conversion of ferrous into ferric salts, and enables the next reagent (ammonia) *completely* to precipitate the iron. Add excess of ammonia, and shake the mixture. Filter.

Precipitate Al Fe*.		Filtrate Zn.
Dissolve in HCl, add excess of KHO, stir, filter.		Test by AmHS (white ppt.).
Ppt. Fe (red ppt.).	Filtrate Al. Make slightly acid by HCl, and add excess of AmHO† (white ppt.).	

Note I.—If iron is present, portions of the original solution must be tested by ferridcyanide of potassium for ferrous, and by ferrocyanide for ferric salts; dark-blue precipitates with both indicate both salts.

Note II.—If no ferrous salt is present, ebullition with nitric

* The aluminium precipitate (Al_2O_3) is white, the iron (Fe_2O_3) red. If the precipitate is red, iron must be and aluminium may be present; if white, iron is absent, and further operations on the ppt. are unnecessary.

This precipitate (Al_2O_3 and Fe_2O_3) may also, if sufficient is at disposal, be analysed by simply well shaking a washed portion in a tube with solution of potash or soda; the hydrate of iron is not thereby affected, while the hydrate of aluminium is dissolved, and may be detected in the clear decanted fluid by neutralizing all alkali by a little excess of acid, and then adding excess of ammonia.

† Alumina, when in small quantity, is sometimes prevented from being precipitated by ammonia through the presence of organic matter derived from the filter-paper by action of the potash. In cases of doubt, therefore, before adding ammonia boil the liquid with a little nitric acid, which destroys any organic matter.

acid is unnecessary. It is, perhaps, therefore advisable always to determine this point by *previously* testing a little of the original solution with ferridcyanide; if no blue precipitate occurs, the nitric-acid treatment may be omitted.

CHART FOR ALL METALS HITHERTO CONSIDERED.

The following Table (*vide* p. 183) is perhaps the best, but not the only adaptation of the ordinary reactions to systematic analysis. In it, the analytical scheme for the third group is added to that of the first two groups. As before, analysis is commenced by the addition of chloride of ammonium to prevent partial precipitation of magnesium, and by ammonia, to neutralize any acid—for acid destroys the group-precipitant, sulphhydrate of ammonium, preventing its useful action and causing a precipitation of the free sulphur it commonly contains. Any precipitate by the ammonia may be disregarded, for the sulphhydrate attacks both solid and liquid.

Note.—When a test gives no reaction, absence of the body sought may fairly be inferred. If a group-test (that is, a test which precipitates a group of substances) gives no reaction, the analyst is saved the trouble of looking for any of the members of that group.

QUESTIONS AND EXERCISES.

206. Name the chief ores of iron.
207. How is the metal obtained from the ores?
208. What is the chemical difference between cast iron, wrought iron, and steel?
209. Explain the process of "welding."
210. What is the nature of chalybeate waters?
211. Illustrate by formulæ the difference between ferrous and ferric salts.
212. Under what different circumstances may the atom of iron be considered to exert bivalent, trivalent, and sexivalent activity?
213. Write a paragraph on the nomenclature of iron salts.
214. Give a diagram of the official process for the preparation of ferrous sulphate.
215. In what respects do the official Sulphate of Iron, Granulated Sulphate of Iron, and dried Sulphate of Iron differ?

216. How is ferrous sulphate obtained on the large scale?

217. Mention the chemical names of white, green, and blue vitriol.

218. Why does ferrous sulphate become brown by prolonged exposure to air?

219. Give a diagram showing the formation of Ferrous Carbonate.

220. Describe the action of atmospheric oxygen on ferrous carbonate; can the effect be prevented?

221. In what order would you mix the ingredients of *Mistura Ferri Composita*, and why?

222. Write out an equation illustrative of the formation of the official Phosphate of Iron.

223. Why is acetate of sodium used in the preparation of ferrous phosphate?

224. Which four compounds of iron may be formed by the direct union of their elements?

225. Give the official method for the preparation of Solution of Ferric Chloride.

226. Of what use is the spirit in Tincture of Perochloride of Iron?

227. How may Ferrous be converted in Ferric Sulphate?

228. What is the formula of Ferric Acetate? and how is it prepared for use in pharmacy?

229. Express, by formulæ, the difference between the *Ferri Peroxidum Humidum*, B. P., and *Ferri Peroxidum Hydratum*, B. P.

230. How does Ferric Hydrate act as an antidote to arsenic?

231. What are the properties of anhydrous ferric oxide?

232. What are the general characters and mode of production of the medicinal scale preparations of iron?

233. In what state is the iron in *Vinum Ferri*, B. P.?

234. What other form of Wine of Iron is official?

235. Give equations illustrating the chief steps in the artificial production of the so-called Magnetic Oxide of Iron.

236. How is precipitated magnetic oxide of iron distinguished from the varieties made directly from the metal?

237. Why is magnetic oxide of iron officially directed to be dried at a temperature not exceeding 120° Fahr.?

238. Give a diagram showing the formation of Ferric Nitrate.

239. Work out a sum showing how much anhydrous ferric oxide will yield, theoretically, one hundredweight of iron. Ans. 160 lbs.

TABLE OF SHORT DIRECTIONS FOR THE ANALYSIS OF AN AQUEOUS SOLUTION OF SALTS OF ANY OR ALL OF THE METALLIC ELEMENTS HITHERTO CONSIDERED.

Add AmCl ; AmHO ; AmHS ; stir, filter.

Precipitate Fe Al Zn. Wash, dissolve in HCl¹, boil (to remove H₂S), filter (to remove S), add excess of KHO¹, stir, filter.		Filtrate Ba Ca Mg Am Na K. Add Am₂CO₃, boil, filter.			
Ppt. Fe (test orig. sol. by K₄FeO₇ and K₂FeO₄).	Filtrate Al Zn. Neutralize with HCl, add AmHO, stir, filter.	Ppt. Ba Ca. Dissolve in HCl, H₂O,¹ add K₂CrO₄, filter.	Filtrate Mg Am Na K. Add Am₂HPO₄, stir, filter.		
Ppt. Al (white).	Filtrate Zn Add AmHS (white ppt.).	Ppt. Ba (yellow).	Filtrate Ca. Add Am₂C₂O₄, (white ppt.).	Ppt. Mg (white).	Filtrate Am Na K. Evap., ignite, dissolve. Na by flame; K by PtCl₄; orig. sol. for Am.

* Add, also, a few drops of HNO_3 if iron be present—i.e. if the ppt. be black. (Vide notes on pp. 177 and 180.)
 † Or add excess of ammonia, filter, and test filtrate for zinc. The hydrates of iron and aluminium may then be separated by an alkali, as described in the footnote to the previous Table (p. 180).

240. Give the characteristic tests for iron, distinguishing between ferrous and ferric reactions, and illustrating each by an equation or a diagram:—

- a. Sulphydrate of ammonium.
- b. Ferrocyanide of potassium.
- c. Ferridcyanide of potassium.
- d. Caustic alkalies.
- e. Sulphocyanate of potassium.

241. Describe the action of ammonia on salts of iron, aluminium, and zinc respectively.

242. What precautions must be used in testing for calcium a solution containing iron?

243. How is magnesium detected in the presence of zinc?

244. How is aluminium detected in presence of magnesium?

245. Draw up a scheme for the analysis of an aqueous liquid containing salts of iron, barium, and potassium.

246. How may zinc, magnesium, and ammonium be consecutively removed from aqueous solution?

ARSENICUM, ANTIMONY.

These elements resemble metals in appearance and in the character of some of their compounds; but they are still more closely allied to the non-metals, especially to phosphorus and nitrogen. Their atoms are quinquivalent (As^V , Sb^V), as seen in arsenic anhydride (As_2O_5) and pentachloride of antimony (SbCl_5), but usually exert trivalent activity only (As^{III} , Sb^{III}), as seen in the hydrogen and other compounds (AsH_3 , AsCl_3 , AsBr_3 , AsI_3). A few preparations of these elements are used in medicine; but all are more or less powerful poisons, and hence have considerable toxicological interest.

Arsenicum is an exception to the rule that the atomic weights (taken in grains, grammes, or other weight) of elements, under similar circumstances of temperature and pressure, give equal volumes of vapour, the equivalent weight (75) of arsenicum only occupying half such a volume. Hence while the molecular weights (that is, double the atomic weights) of oxygen ($\text{O}_2 = 32$), hydrogen ($\text{H}_2 = 2$), nitrogen ($\text{N}_2 = 28$), etc. give a similar bulk of vapour at any given temperature and pressure, the double atomic weight of arsenicum ($\text{As}_2 = 150$), at the same temperature and pressure, only

affords half this bulk. It would appear, therefore, that the molecule of arsenicum contains four atoms, and that its formula is As_4 . As in the case of sulphur, however, arsenicum, in the state ordinarily known to us, may be abnormal, and a variety yet be found in which the molecular weight is double (instead of quadruple) the atomic weight.

From observed analogy between the two metals, the molecular constitution of antimony is probably similar to that of arsenicum.

ARSENICUM.

Symbol As . Atomic weight 75.

Sources.—Arsenical ores are frequently met with in nature, the commonest being the arsenio-sulphide of iron ($FeSAs$). This mineral is roasted in a current of air, the oxygen of which, combining with the arsenicum, forms common white arsenic (As_2O_3 or, perhaps, As_4O_6), (*Acidum Arseniosum*, B. P.), which is condensed in chambers or long flues. It commonly "occurs as a heavy white powder, or in sublimed masses, which usually present a stratified appearance, caused by the existence of separate layers, differing from each other in degrees of opacity." The vitreous or amorphous arsenic is far more soluble than the crystalline variety, and in other respects they differ in properties. Such differences between the crystalline and amorphous varieties of an element or compound are not unfrequent: they have not yet been satisfactorily explained. *Realgar* (red algar) is the red native sulphide (As_2S_2), and orpiment (*auripigmentum*, the golden pigment), the yellow native sulphide (As_2S_3) of arsenicum. The iodide of arsenicum (AsI_3) may be made from its elements or (Babcock) by dissolving white arsenic in aqueous hydriodic acid and evaporating.

REACTIONS HAVING (a) SYNTHETICAL AND (b) ANALYTICAL INTEREST.

(a) *Reactions having Synthetical Interest.*

Alkaline Solution of Arsenic.

First Synthetical Reaction.—Boil a grain or two of powdered arsenic (As_2O_3) in water containing a little carbonate of potassium, and, if necessary, filter. The solution, coloured with

compound tincture of lavender, and containing 4 grains of arsenic per ounce, forms the *Liquor Arsenicalis*, B. P. (*Fowler's Solution*).

Note.—This official solution does not generally contain arsenite of potassium; for the arsenic does not decompose the carbonate of potassium, or only after long boiling. From concentrated solutions carbonic acid gas is more quickly eliminated.

Arsenious Acid and other Arsenites.

Arsenic, or *arsenious anhydride* (the so-called arsenious acid), when dissolved in water is said to yield true arsenious acid (H_3AsO_3), the arsenite of hydrogen.



When arsenic is dissolved in excess of solutions of potash or soda, arsenites are formed having the formulæ KH_2AsO_3 and NaH_2AsO_3 . Boiled with excess of arsenic one molecule of these salts combines with one of the arsenic. The usual character of such compounds is that of oily alkaline liquids.

Arsenic fused with alkaline carbonates yields pyroarsenites ($\text{Na}_4\text{As}_2\text{O}_7$ or $\text{K}_4\text{As}_2\text{O}_7$) and metallic arsenicum. Arsenites have the general formula $\text{E}'_3\text{AsO}_3$.

Acid Solution of Arsenic.

Second Synthetical Reaction.—Boil arsenic with dilute hydrochloric acid. Such a solution made with prescribed proportions of acid and water, and containing 4 grains of arsenic (As_2O_3) per ounce, forms the *Liquor Arsenici Hydrochloricus*, B. P. (*De Valangin's Solution* contained a grain and a half per ounce.)

Note.—No decomposition occurs in this experiment. The liquid is simply a solution of arsenic in dilute hydrochloric acid. These two solutions may be preserved for analytical operations.

Mem.—The practical student should boil arsenic in water only, and thus have an acid, alkaline, and aqueous solution for analytical comparison.

Arsenicum.

Third Synthetical Reaction.—Place a grain or less of arsenic at the bottom of a narrow test-tube, cover it with about half an inch or an inch of small fragments of dry charcoal, and hold the tube, nearly horizontally, in a flame, the mouth being loosely covered by the thumb. At first let the bottom of the tube project slightly beyond the flame, so that the charcoal may become nearly red-hot; then heat the bottom of the tube. The arsenic will sublime, become deoxidized by the charcoal, carbonic oxide being formed, and arsenicum be deposited in the cooler part of the tube as a dark mirror-like metallic incrustation.

There is a characteristic odour, resembling garlic, emitted during this operation, probably due to a partially oxidized trace of arsenicum which escapes from the tube; for arsenic does not give this odour; moreover, arsenicum being a freely oxidizable element, its vaporous particles could scarcely exist in the air in an entirely unoxidized state.

Metallic arsenicum may be obtained in large quantities by the above process if the operation be conducted in vessels of commensurate size. But performed with great care, in narrow tubes, using not charcoal alone, but *black flux* (a mixture of charcoal and carbonate of potassium obtained by heating acid tartrate of potassium in a test-tube or other closed vessel till no more fumes are evolved), the reaction has considerable analytical interest, the garlic odour and the formation of the mirror-like ring being highly characteristic of arsenicum. Compounds of mercury and antimony, however, give sublimes which may be mistaken for arsenicum.

Arsenic Acid and other Arseniates.

Fourth Synthetical Reaction.—Boil a grain or two of arsenic with a few drops of nitric acid until red fumes cease to be evolved; evaporate the solution in a small dish to dryness, to remove excess of nitric acid; dissolve the residue in water: the product is Arsenic acid (H_3AsO_4).

Arsenic acid, when strongly heated, loses the elements of water, and arsenic anhydride remains (As_2O_5).

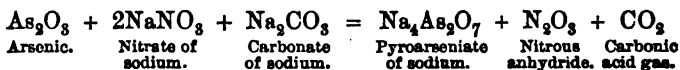
Arsenic anhydride readily absorbs water and becomes arsenic acid (H_3AsO_4). Arsenic acid is easily reduced to arsenious by the action of reducing agents, such as sulphurous acid ($\text{H}_3\text{AsO}_4 + \text{H}_2\text{SO}_3 = \text{H}_3\text{AsO}_3 + \text{H}_2\text{SO}_4$) or oxalic acid ($\text{H}_3\text{AsO}_4 + \text{H}_2\text{C}_2\text{O}_4 = \text{H}_3\text{AsO}_3 + \text{H}_2\text{O} + 2\text{CO}_2$).

Salts analogous to arsenic acid, the arseniate of hydrogen, are termed *arseniates*, and have the general formula $\text{R}'_3\text{AsO}_4$. The ammonium arseniate (Am_2HAsO_4) may be made by neutralizing arsenic acid with ammonia. Its solution in water forms a useful reagent.—Arsenic acid is used as an oxidizing agent in the manufacture of the well-known dye, magenta.

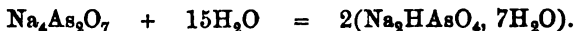
Arsenite and arseniate of sodium are used in the cleansing-operations of the calico-printer.

Pyroarsenate and Arseniate of Sodium.

Fifth Synthetical Reaction.—Fuse two or three grains of common white arsenic (As_2O_3) with nitrate of sodium (NaNO_3) and dried carbonate of sodium (Na_2CO_3) in a porcelain crucible, and dissolve the mass in water; solution of arseniate of sodium (Na_2HAsO_4) results.



The official proportions are 10 of arsenic to $8\frac{1}{2}$ of nitrate of sodium and $5\frac{1}{2}$ of dried carbonate, each powdered, the whole well mixed, fused in a crucible at a red heat till effervescence ceases, and the liquid poured out on a slab. The product is pyroarsenate of sodium ($\text{Na}_4\text{As}_2\text{O}_7$). Dissolved in water, crystallized, and dried, the salt has the formula $\text{Na}_2\text{HAsO}_4 \cdot 7\text{H}_2\text{O}$ (*Sodæ Arsenias*, B. P.; *Sodii Arsenias*, U. S. P.).



Heated to 300°F . the crystals lose all water. A solution of 4 grains of the anhydrous salt (Na_2HAsO_4) in 1 ounce of water (equivalent to about 2 grains of arsenic, and therefore about half the strength of *Liquor Arsenicalis*, B. P.) forms the *Liquor Sodæ Arseniatis*, B. P. The anhydrous salt is used in this preparation because the crystallized is of somewhat uncertain composition. The fresh crystals are represented by

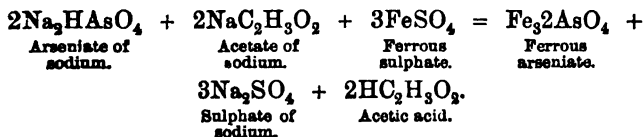
the formula $\text{Na}_2\text{HAsO}_4, 12\text{H}_2\text{O}$ (= 53.7 per cent. of water); these soon effloresce and yield a stable salt having the formula $\text{Na}_2\text{HAsO}_4, 7\text{H}_2\text{O}$ (= 40.4 per cent. of water). To avoid the possible employment of a mixture of these bodies, the invariable anhydrous salt is officially used, constancy in the strength of a powerful preparation being thereby secured.

The student will find useful practice in verifying, by calculation, the above numbers representing the centesimal proportion of water in the two arseniates of sodium. This will be easy if what has already been stated respecting a symbol representing a number as well as a name, and the remarks concerning molecular weight, be remembered.

The shape of each of the two varieties of arseniate of sodium ($\text{Na}_2\text{HAsO}_4, 12\text{H}_2\text{O}$, and $\text{Na}_2\text{HAsO}_4, 7\text{H}_2\text{O}$) is identical with that of the corresponding phosphate of sodium ($\text{Na}_2\text{HPO}_4, 12\text{H}_2\text{O}$, and $\text{Na}_2\text{HPO}_4, 7\text{H}_2\text{O}$); the structure of the molecule of the 12-arseniate is the same as that of the 12-phosphate, and the 7-arseniate as that of the 7-phosphate; the two former are isomorphous, the two latter are isomorphous. This is only one instance of the strong analogy of arsenicum and its compounds with phosphorus and its corresponding compounds. The preparation and characters of the next substance, arseniate of iron, will remind the learner of phosphate of iron.

Arseniate of Iron. Ferrous Arseniate.

Sixth Synthetical Reaction.—To solution of arseniate of sodium add a little acetate of sodium and then solution of ferrous sulphate; a precipitate of ferrous arseniate occurs (Fe_32AsO_4) (*Ferri Arsenias*, B. P.). On the large scale 4 parts of dried arseniate and 3 of acetate dissolved in 40 of water, mixed with 9 of sulphate in 60 of water, may be employed. The precipitate should be collected on a calico filter, washed, squeezed, and dried at a low temperature (100°F .) over a water-bath to avoid excessive oxidation.



The use of the acetate of sodium is to insure the absence of free sulphuric acid in the solution—sulphate of sodium being formed together with acetic acid. Sulphuric acid is a solvent of ferrous arseniate; acetic acid is not. It is impossible to prevent the separation of sulphuric acid, if only ferrous sulphate and arseniate of sodium be employed.

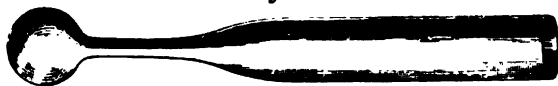
At the instant of precipitation ferrous arseniate is white, but rapidly becomes of a green or greenish-blue colour owing to absorption of oxygen and formation of a ferroso-ferric arseniate. It is a tasteless amorphous powder, soluble in acids.

The Hydride and Sulphides of Arsenicum, and the Arsenites and Arseniates of Copper and of Silver, are mentioned in the following analytical paragraphs.

(b) *Reactions having Analytical Interest (Tests).*

First Analytical Reaction.—Repeat the third synthetical reaction, operating on not more arsenic than has about the bulk of a small pin's head, and using not charcoal alone, but the *black flux* already mentioned (p. 187) or a well made and perfectly dry mixture of charcoal and carbonate of potassium, the latter best obtained by heating bicarbonate of potassium. The tube employed should be a narrow test-tube, or, better, a tube (easily made from glass tubing) having more or less the following (Berzelius's) form :—

Fig. 33.



The arsenic and black flux are placed in the bulb of the tube, which is then heated in a flame; the arsenicum condenses on the constricted portion of the tube. If now the bulb be carefully fused off in a flame, the arsenicum may be chased up and down the narrower part of the tube until the air in the tube has re-oxidized it to arsenious anhydride.

If the operation has been performed in a less delicate

manner in an ordinary test-tube, cut or break off portions of the tube containing the sublimate of arsenicum, put them into a test-tube and heat the bottom of the latter, holding it nearly horizontally, and partially covering the mouth with the finger or thumb; the arsenicum (As_4) will absorb oxygen from the air in the tube, and the resulting arsenious anhydride (As_2O_3) be deposited on the cool part of the tube in brilliant transparent, generally imperfect, octahedral crystals.

Fig. 34.



A SUBLIMATE OF ARSENIC (MAGNIFIED).

Fig. 34a.



A PERFECT OCTAHEDRON.

Microscopic Test.—Prove that the crystals are identical in form with those of common white arsenic, by heating a grain or less of the latter in another test-tube, examining the two sublimates by a good lens or compound microscope.

The appearance of a sublimate of arsenic is peculiar and quite characteristic. The primary form of each crystal is an octahedron (*ὀκτώ*, *okto*, eight; *ἑδρα*, *hedra*, side) (fig. 34a), or, rarely, a tetrahedron, and in a sublimate a few perfect octahedra are generally present. Usually, however, the crystals are modifications of octahedra such as are shown in fig. 34—which is drawn from actual sublimates.

Second Analytical Reaction.—Place a thin piece of copper, about a quarter-inch wide and half-inch long, in a solution of arsenic, acidified by hydrochloric acid, and boil (nitric acid must not be present, or the piece of metal will be dissolved); arsenicum is deposited on the copper in a metallic condition, an equivalent portion of copper going into solution. Pour off

the supernatant liquid from the copper, wash the latter once or twice with water, dry the piece of metal by holding in the fingers and passing through a flame, and finally place it at the bottom of a clean dry narrow test-tube or a Berzelius tube and sublime as described in the last reaction, again noticing the form of the resulting crystals.

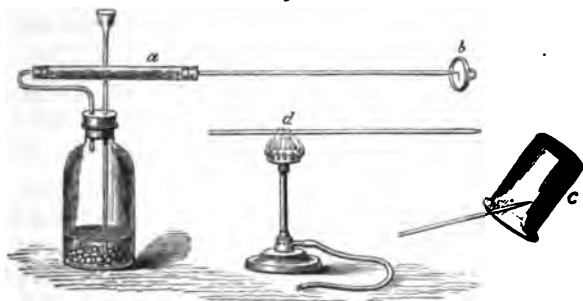
This is commonly known as Reinsch's test for arsenicum. The tube may be reserved for subsequent comparison with an antimonial sublimate.

Note.—Copper itself frequently contains arsenicum, a fact that may not, perhaps, much trouble an operator so long as he is performing experiments in practical chemistry merely for educational purposes; but when he engages in the analysis of bodies of unknown composition, he must assure himself that neither his apparatus nor materials already contain the element for which he is in search.

The detection of arsenicum in metallic copper is best accomplished by distilling a mixture of a few grains of the sample with five or six times its weight of ferric hydrate or chloride (free from arsenicum) and excess of hydrochloric acid. The arsenicum is thus volatilized in the form of chloride of arsenicum, and may be condensed in water and detected by sulphuretted hydrogen (6th Analytical Reaction) or Reinsch's test. The ferric chloride solution is, if necessary, freed from any trace of arsenicum by evaporating once or twice to dryness with excess of hydrochloric acid.

Third Analytical Reaction—The hydrogen test, or "Marsh's" test.—Generate hydrogen in the usual way from water by zinc and sulphuric acid, a bottle of about four or six ounces capacity being used, and a funnel-tube and short delivery-tube passing through the cork in the usual manner (see following figure). Dry the escaping hydrogen (except in rough experiments, when it is unnecessary) by adapting to the delivery-tube, by a pierced cork, a short piece of wider tubing filled with fragments of chloride of calcium (a). To the opposite end of the drying-tube fit a piece of narrow tubing ten or twelve inches long, made of hard German glass, and having its aperture narrowed by drawing out in the flame of the blow-pipe.

Fig. 35.



THE HYDROGEN TEST FOR ARSENICUM.

When the hydrogen has been escaping for a sufficient number of minutes and at such a rate as to warrant the operator in concluding that all the air originally existing in the bottle has been expelled, set light to the jet, and then pour eight or ten drops of the aqueous solution of arsenic, or three or four drops of the acid or alkaline solution of arsenic, previously prepared, into the funnel-tube, washing the liquid into the generating-bottle with a little water. The arsenic is at once reduced to the state of arsenicum, and the latter combines with some of the hydrogen to form hydride of arsenicum or arseniuretted hydrogen gas (AsH_3). Immediately hold a piece of earthenware or porcelain (the lid of a porcelain crucible (*b*), if at hand) in the hydrogen jet at the extremity of the delivery tube; a brown spot of arsenicum is deposited on the porcelain. Collect several of these spots, and retain them for future comparison with antimonial spots similarly obtained (p. 207).

The separation of arsenicum in the flame is due to the decomposition of the arseniuretted hydrogen by the heat of combustion. The cool porcelain at once condenses the arsenicum, and thus prevents its oxidation to white arsenic, which would otherwise take place at the outer edge of the flame.

Hold a small beaker (*c*) or wide test-tube over the flame for a few minutes; a white film of arsenic (As_2O_3) will be

slowly deposited, and may be further examined in contrast with a similar antimonial film obtained in the same manner (p. 207).

During these experiments the effect produced by the arsenical vapours on the colour of the hydrogen-flame will have been noticed; they give it a dull livid bluish tint. This is characteristic.

Apply the flame of a gas-lamp to the middle of the hard glass delivery-tube *d*; the arseniuretted hydrogen, as before, is decomposed by the heat, but the liberated arsenicum (As_4) immediately condenses in the cool part of the tube beyond the flame, forming a dark metallic mirror. The tube may be removed and kept for comparison with an antimonial deposit (p. 207).

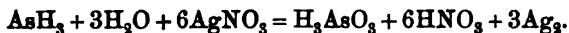
Note I.—Zinc, like copper, frequently itself contains arsenicum. When a specimen free from arsenicum is met with, it should be reserved for analytical experiments, or a quantity of guaranteed purity should be purchased of the chemical-apparatus maker. Sulphuric acid is more easily obtained free from arsenic.

Note II.—In delicate and important applications of Marsh's test, magnesium may be substituted for zinc with safety, as arsenicum has not yet been, and is not likely to be, found in magnesium. Magnesium in rods is convenient for this purpose, and may be obtained from most dealers in chemicals.

Note III.—Sulphuric acid, which is often used for drying gases, decomposes arseniuretted hydrogen. Chloride of calcium is therefore the appropriate desiccating agent for this gas.

Fourth Analytical Reaction—Fleitmann's test.—Generate hydrogen by heating in a test-tube to near the boiling-point a strong solution of caustic soda or potash and some pieces of zinc ($Zn + 2NaHO = H_2 + Na_2ZnO_2$ —zincate of sodium). Add a drop of arsenical solution. Now spread over the mouth of the tube a cap of filter-paper moistened with one drop of solution of nitrate of silver. Again heat the tube, taking care that the liquid itself shall not spurt up on to the cap. A plug of cotton wool may even be placed in the mouth of the

test-tube to prevent this spurting (Senier). The arsenic is reduced to arsenicium, the latter uniting with the hydrogen as in Marsh's test; and the arseniuretted hydrogen passing up through the cap reacts on the nitrate of silver, causing the production of a purplish-black spot.



Note I.—This reaction is particularly valuable, enabling the analyst to quickly distinguish arsenicium in the presence of its sister element antimony, which, although it combines with the hydrogen evolved from dilute acid and zinc, does not combine with the hydrogen evolved from solution of alkali and zinc, and therefore does not give the effect just described.

Note II.—Aluminium answers as well as zinc for Fleitmann's test (Gatehouse), or magnesium may be used; or instead of zinc and alkali weak sodium amalgam may be employed (Davy).

Fifth Analytical Reaction.—To a solution of chloride of tin in strong hydrochloric acid add a very small quantity of any arsenical solution. Arsenicium then separates, especially on the application of heat, giving the mixture a yellowish and then brownish hue or greyish-brown turbidity, or even a sediment of grey-brown flocks, according to the amount present. Much water prevents the reaction; its presence therefore must be avoided as far as possible; indeed a liquid saturated by hydrochloric acid gas gives best results. Arsenic in sulphuric or hydrochloric acid or in tartar emetic, etc., may be detected by this method. Nitrates, such as subnitrate of bismuth, must first be heated with sulphuric acid to remove the nitric radical before applying this reduction test for arsenicium. The stannous is converted to stannic salt during the reaction.

Distinction between Arsenious and Arsenic combinations.—The above tests are those of arsenicium, whether existing in the arsenious or arsenic condition, though from the latter the element is not generally eliminated so quickly as from the former. Of the following reactions, that with nitrate of silver

at once distinguishes arsenious acid and other arsenites from arsenic acid and other arseniates.

Mem.—The exact nature of all these analytical reactions will be more fully evident if traced out by diagrams or equations.

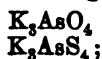
Sixth Analytical Reaction.—Through an acidified solution of arsenic pass sulphuretted hydrogen; a yellow precipitate of sulphide of arsenicum or arsenious sulphide (As_2S_3) quickly falls. Add an alkaline hydrate or sulphhydrate to a portion of the precipitate, it readily dissolves. The precipitate consequently would not be obtained on passing sulphuretted hydrogen through an alkaline solution of arsenic. To another portion of the precipitate, well drained, add *strong* hydrochloric acid; it is insoluble—unlike sulphide of antimony. (Neither sulphide is soluble in the *weak* acid.)

Note I.—Cadmium also affords a yellow sulphide in an acid solution by action of sulphuretted hydrogen; but this sulphide is insoluble in alkaline liquids. Under certain circumstances tin, too, yields a yellow sulphide, but tin is otherwise easily distinguished (*vide* "Tin" in Index).

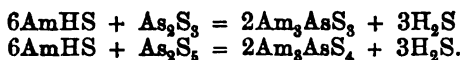
Note II.—A trace of sulphide of arsenicum is sometimes met with in sulphur (distilled from arsenical pyrites). It may be detected by digesting the sulphur in solution of ammonia, filtering, and evaporating to dryness; a yellow residue of sulphide of arsenicum is obtained if that substance be present.

Seventh Analytical Reaction.—Through an acidified solution of arsenic acid, or any other arseniate, pass sulphuretted hydrogen; the arsenic compound is gradually reduced to the arsenious and a yellow precipitate of arsenious sulphide and sulphur ($\text{As}_2\text{S}_3 + \text{S}_2$) slowly falls, soluble in alkaline hydrates and sulphhydrates.

Chemical Analogy of Sulphur and Oxygen.—The potassium arsenite and sulph-arsenite, arseniate and sulph-arseniate, have the composition represented by the following formulæ:—



and the corresponding ammonium and sodium salts have a similar composition :—



Eighth Analytical Reaction.—To an aqueous solution of arsenic add two or three drops of solution of sulphate of copper, and then cautiously add diluted solution of ammonia, drop by drop, until a green precipitate is obtained. The production of this precipitate is characteristic of arsenicum. To a portion of the mixture add an acid; the precipitate dissolves. To another portion add alkali; the precipitate dissolves. These two experiments show the advantage of testing a suspected arsenical solution by litmus-paper before applying this reaction—if acid, cautiously adding alkali, if alkaline, adding acid, till neutrality is obtained. Or a special copper reagent may be used: see a note to the Eleventh Analytical Reaction.

The precipitate is arsenite of copper ($\text{Cu}^{\text{H}}\text{HASO}_3$) or *Scheele's Green*. More or less pure, or mixed with acetate or, occasionally, carbonate of copper, it is very largely used as a pigment under many names, such as Brunswick Green and Schweinfurth Green, by painters, paper-stainers, and others.

Ninth Analytical Reaction.—Apply the test just described to a solution of arsenic acid or other arseniate; a somewhat similar precipitate of arseniate of copper is obtained.

Tenth Analytical Reaction.—Repeat the eighth reaction, substituting nitrate of silver for sulphate of copper: in this case *yellow* arsenite of silver (Ag_3AsO_3) falls, also soluble in acids and alkalies.

Eleventh Analytical Reaction.—Apply the silver test to a solution of arsenic acid or other arseniate; a *chocolate-coloured* precipitate of arseniate of silver (Ag_3AsO_4) falls.

Copper and Silver Reagents for Arsenicum.—The last four reactions may be performed with increased delicacy and certainty of result if the copper and silver reagents be previously prepared in the following manner:—To solution

of pure sulphate of copper (about 1 part in 20 of water) add ammonia until the blue precipitate at first formed is nearly, but not quite, redissolved; filter and preserve the liquid as an arsenicum reagent, labelling it *solution of ammonio-sulphate of copper* (B. P.). Treat solution of nitrate of silver (about 1 part in 40) in the same way, and label it *solution of ammonio-nitrate of silver* (B. P.). The composition of these two salts will be referred to subsequently.

Arsenious and Arsenic Compounds.—While many reagents may be used for the detection of arsenicum, only nitrate of silver, as already stated, will readily indicate in which state of oxidation the arsenicum exists; for the two sulphides and the two copper precipitates, though differing in composition, resemble each other in appearance, whereas the two silver precipitates differ in colour as well as in composition.

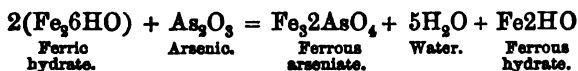
Soluble arseniates also give insoluble arseniates with solutions of salts of barium, calcium, zinc, and some other metals.

In group-testing, arsenicum, if existing as arsenic acid or other arseniate, is not readily affected by such tests as sulphuretted hydrogen or even hydrogen itself. Hence, if its presence in that state is suspected, the liquid under analysis should be warmed with a little sulphurous acid or oxalic acid (*vide* p. 188) and then tested with sulphuretted hydrogen.

Antidote.—In cases of poisoning by arsenic or arsenical preparations, the most effective antidote is recently precipitated moist ferric hydrate (*Ferri Peroxidum Humidum*, B. P.). It is perhaps best administered in the form of a mixture of solution of perchloride of iron (*Liquor Ferri Perchloridi*, B. P.) with carbonate of sodium—two to three ounces of the former to about one ounce of the crystals of the latter. Instead of the carbonate of sodium about a quarter of an ounce of calcined magnesia (*Magnesia*, B. P.) may be used. These quantities will render at least 10 grains of arsenic insoluble. Emetics should also be given, and the stomach-pump applied as quickly as possible.

The above statements regarding the antidote for arsenic may be verified by mixing the various substances together, filtering, and proving the absence of arsenicum in the filtrate by applying some of the foregoing tests.

Mode of action of the Antidote.—The action of the carbonate of sodium or the magnesia is to precipitate ferric hydrate ($\text{Fe}_2\text{O}_3\cdot 3\text{H}_2\text{O}$)—chloride of sodium (NaCl) or of magnesium (MgCl_2) being formed, which are harmless, if not beneficial, under the circumstances. The reaction between the ferric hydrate and the arsenic results in the formation of insoluble ferrous arseniate.



As already stated, dried ferric hydrate (then become an oxyhydrate, $\text{Fe}_2\text{O}_3\cdot 2\text{H}_2\text{O}$) (*Ferri Peroxidum Hydratum*, B. P.) has less action on arsenic; and even the moist recently prepared hydrate ($\text{Fe}_2\text{O}_3\cdot 3\text{H}_2\text{O}$) loses much of this power as soon as it has become converted into an oxyhydrate ($\text{Fe}_2\text{O}_3\cdot 2\text{H}_2\text{O}$), a change which occurs though the hydrate be kept under water (Procter). According to T. and H. Smith this decomposition occurs gradually, but in an increasing ratio; so that after four months the power of the moist mass is reduced to one-half, and after five months to one-fourth.

The so-called Solution of Dialysed Iron (*see* Index) is also, as might be expected from its composition, an antidote to arsenic. It should be administered with a little bicarbonate of either sodium or potassium, or magnesia, or any other salt which serves to neutralize any acid that may be present.

QUESTIONS AND EXERCISES.

247. What is the formula of a molecule of arsenicum?
248. In what form does arsenicum occur in nature?
249. Describe the characters of white arsenic?
250. Name the official preparations of arsenicum.
251. What proportion of arsenic (As_2O_3) is contained in *Liquor Arsenicalis*, B. P., and *Liquor Arsenici Hydrochloricus*, B. P.?
252. By what method may arsenic be reduced to arsenicum?
253. Give the formulæ of arsenious and arsenic acids and anhydrides.
254. Explain, by diagrams, the reactions which occur in converting arsenic into Arseniate of Sodium by the process of the Pharmacopœia.

255. Why is anhydrous instead of crystallized arseniate of sodium employed in the preparation of *Liquor Sodæ Arseniatis*, B. P.?

256. In the preparation of Arseniate of Iron from ferrous sulphate and arseniate of sodium, why is acetate of sodium included?

257. Describe the manipulations necessary to obtain arsenic in its characteristic crystalline form.

258. How is Reinsch's test for arsenicum applied, and under what circumstances may its indications be fallacious?

259. Give the details of Marsh's test for arsenicum, and the precautions to be observed in its performance. Explain the reactions by diagrams.

260. What peculiar value has Fleitmann's test for arsenic?

261. Describe the conditions under which sulphuretted hydrogen becomes a trustworthy test for arsenicum.

262. How may a trace of sulphide of arsenicum be detected in sulphur?

263. How are salts of copper and silver applied as reagents for the detection of arsenicum?

264. How are arsenites distinguished from arseniates?

265. Mention the best antidote in cases of poisoning by arsenic, explain the process by which it may be most quickly prepared, and describe its action.

266. Do you know of any other antidote to arsenic? if so, describe the mode of administration.

ANTIMONY.

Symbol Sb (stibium). Atomic weight 122.

Source and Uses.—Antimony occurs in nature chiefly as sulphide, Sb_2S_3 . The *crude* or *black antimony* of pharmacy is this native sulphide freed from impurities by fusion; it has a striated, crystalline, lustrous fracture; subsequently powdered it forms the greyish-black crystalline *Antimonium nigrum*, B. P. The metal is obtained from the sulphide by roasting, the resulting oxide being reduced with charcoal and carbonate of sodium. Metallic antimony is an important constituent of *Type-metal*, *Britannia metal* (tea- and coffee-pots, spoons, etc.), and the best varieties of *Pewter*. The old *pocula emetica*, or everlasting emetic cups, were made of antimony;

wine kept in them for a day or two acquired a variable amount of emetic quality. The metal is not used in making the antimonial preparations of the Pharmacopœia, the sulphide alone being, directly or indirectly, employed for this purpose.

Antimony has very close chemical analogies with arsenicum. Its atom, in the common salts, exerts trivalent activity (e.g. SbCl_3), but sometimes it is quinivalent (e.g. SbCl_5).

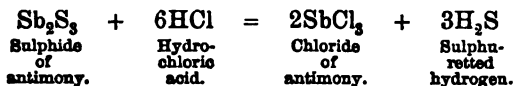
Antimony, like arsenicum, unites with iodine to form a tri-iodide (SbI_3). A bromide (SbBr_3) is also known.

REACTIONS HAVING (a) SYNTHETICAL AND (b) ANALYTICAL INTEREST.

(a) *Reactions having Synthetical Interest.*

Chloride of Antimony. Antimonious Chloride.

First Synthetical Reaction.—Boil half an ounce or less of sulphide of antimony with four or five times its weight of hydrochloric acid in a dish in a fume-chamber or in the open air; sulphuretted hydrogen is evolved and solution of chloride of antimony, SbCl_3 , is obtained.



This solution, cleared by subsidence, is what is commonly known as *Butter of antimony* (*Liquor Antimonii Chloridi*, B.P.). pure sulphide has been used in its preparation the liquid is nearly colourless; but much of that met with in veterinary Pharmacy is simply a by-product in the generation of sulphuretted hydrogen from native sulphide of antimony and hydrochloric acid, and is more or less brown from the presence of chloride of iron. It not unfrequently darkens in colour on keeping; this is due to absorption of oxygen from the air, and conversion of light-coloured ferrous into dark-brown ferric chloride or oxychloride.

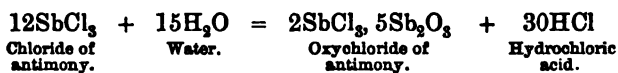
True butter of antimony (SbCl_3) is obtained on evaporating the above solution to a low bulk, and distilling the residue. The butter condenses as a white crystalline semitransparent mass in the neck of the retort; at the close of the operation it may be easily melted and run down into a bottle, which should be subsequently well stoppered.

Pentachloride of antimony (SbCl_5), or *antimonic chloride*, is a fuming liquid, obtained on passing chlorine over the lower chloride.

Oxychloride of Antimony. Antimonious Oxychloride.

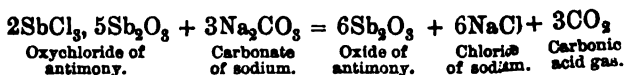
Second Synthetical Reaction.—Pour the solution of chloride of antimony produced in the last reaction into several ounces of water; a white precipitate of oxychloride of antimony ($2\text{SbCl}_3, 5\text{Sb}_2\text{O}_3$) falls, some chloride of antimony remaining in the supernatant acid liquid.

This precipitate is the old *pulvis Algarothi, pulvis angelicus*, or *mercurius vitæ*. It varies somewhat in composition, according to the amount of water with which the chloride may be mixed; but on standing under water gradually becomes crystalline and has the composition above given.



Oxide of Antimony. Antimonious Oxide.

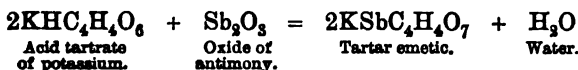
Well wash the precipitate with water, by decantation (*vide* p. 118), and add solution of carbonate of sodium; the chloride remaining with the oxide is thus decomposed, and oxide of antimony (Sb_2O_3) alone remains. This is *Antimonii Oxidum*, B. P. It is of a light buff or greyish-white colour, or quite white if absolutely free from iron, insoluble in water, soluble in hydrochloric acid, fusible at a low red heat. The moist oxide of antimony may be well washed and employed for the next reaction, or dried over a water-bath. At temperatures above 212°F . oxygen is absorbed, and other oxides of antimony formed. The presence of the latter is detected on boiling the powder in solution of acid tartrate of potassium, in which oxide of antimony (Sb_2O_3) is soluble, but *antimonic anhydride* (Sb_2O_5) and the double oxide or so-called *antimonious anhydride* (Sb_4O_8) insoluble.



The higher oxide of antimony (Sb_2O_5), termed antimonie oxide or anhydride, corresponding with arsenic anhydride, is obtained on decomposing the pentachloride by water, or on boiling metallic antimony with nitric acid. The variety obtained from the chloride differs in saturating-power from that obtained from the metal, and is termed metantimonie acid (*μετά, meta, beyond*).

Tartar Emetic.

Third Synthetical Reaction.—Mix the moist oxide of antimony obtained in the previous reaction with about an equal quantity of cream of tartar (6 of the latter to 5 of the dry oxide) and sufficient water to form a paste; set aside for a day to facilitate complete combination; boil the product with water, and filter; the resulting liquid contains the double tartrate of antimony and potassium ($\text{KSbC}_4\text{H}_4\text{O}_7$), potassium-tartrate of antimony, tartarated antimony, or tartar emetic (emetic, from *ἐμέω, emeo*, I vomit; tartar, from *Τάρταρος, tartaros*, see Index).



On evaporation, the salt is obtained in colourless transparent triangular-faced crystals of the above composition, with a molecule of water of crystallization, forming the *Antimonium Tartaratum*, B. P., *Antimonii et Potassii Tartras*, U. S. P. ($\text{KSbC}_4\text{H}_4\text{O}_7, \text{H}_2\text{O}$).

The formula for tartar emetic is apparently inconsistent with the general formula for tartrates ($\text{R}'\text{R}''\text{C}_4\text{H}_4\text{O}_6$); this will be subsequently fully explained in connection with Tartaric Acid. The salt appears to be an oxytartrate $\text{K}'\text{Sb}''(\text{C}_4\text{H}_4\text{O}_6)'''\text{O}''$.

Tartar emetic is soluble in water, and slightly so in proof spirit. Dissolved in sherry wine it forms the official *Vinum Antimoniale*, B. P. It may be externally applied as an ointment, *Unguentum Antimonii Tartarati*, B. P.

Sulphurated Antimony. Oxysulphide of Antimony.

Fourth Synthetical Reaction.—Boil a few grains of sulphide of antimony with solution of soda in a test-tube, and filter; (or larger quantities in larger vessels, 10 ounces of sulphide to $4\frac{1}{2}$ pints of the official solution of soda for 2 hours, frequently stirring, and occasionally replacing water lost by evaporation). Into the filtrate, before cool, stir diluted sulphuric acid until the liquid is slightly acid to test-paper; a brownish-red precipitate of oxysulphide of antimony, the *Antimonium Sulphuratum*, B. P., falls; filter, wash, and dry over a water-bath. It is a mixture of sulphide of antimony (Sb_2S_3) with a small and variable amount of oxide (Sb_2O_3). The oxide results from the double decomposition of sulphide of antimony and soda.

If a small quantity of sulphur be boiled with the sulphide of antimony and solution of soda, the precipitate on addition of sulphuric acid will have a much brighter colour, chiefly on account of the presence of a higher sulphide having a yellow colour (Sb_2S_5).

This is one of the many varieties of *mineral kermes*, so called from their similarity in colour to the *insect kermes*. Kermes is the name, now obsolete, of the *Coccus Ilicis*, a sort of cochineal-insect, full of reddish juice, and used for dyeing from the earliest times. The colour of the precipitate is affected by the temperature as well as state of dilution of the alkaline liquid when the acid is added. When the alkaline liquid is boiled, especially if long exposed to air, oxygen is absorbed by some of the antimony, whose sulphur uniting with the trisulphide forms a portion of the lighter yellow pentasulphide. *Kermes mineral* thus varies much in proportion of oxide and of pentasulphide, also in the physical condition of its trisulphide.

Explanation of process.—The sulphides and oxides of antimony, like those of arsenicum, react with the sulphides and oxides of certain metals to form soluble salts (Na_3SbS_3 and Na_3SbO_3 : the former is deposited in yellow tetrahedral crystals when such an alkaline solution is set aside slowly to cool).

(b) *Reactions having Analytical Interest (Tests).*

First Analytical Reaction.—Through an acidified antimonial solution pass sulphuretted hydrogen; an orange precipitate of amorphous sulphide of antimony falls. It has the same composition as the crystalline black sulphide (Sb_2S_3), into which, indeed, when dried, it is quickly converted by heat. Like sulphide of arsenicum, it is soluble in alkaline solutions. Collect a portion on a filter and, when well drained, add *strong* hydrochloric acid; it dissolves—unlike sulphide of arsenicum.

A higher sulphide of antimony (Sb_2S_5), corresponding to the higher sulphide of arsenicum, exists. It is formed on passing sulphuretted hydrogen through an acidified solution of the higher chloride (SbCl_5), or, less pure, on boiling black sulphide of antimony and sulphur with an alkali, and decomposing the resulting filtered liquid by an acid.

Note.—The arsenious and antimonious compounds are those chiefly employed in medicine; arseniates of sodium and iron are, however, sometimes employed. The arseniates and, rarely, an antimoniate are useful in analysis, and the antimonie chloride in chemical research. The higher compounds of both elements are noticed here chiefly to draw attention to the close analogy existing between arsenicum and antimony, an analogy carried out in the numerous other compounds of these elements.

Second Analytical Reaction.—Dilute two or three drops of the solution of chloride of antimony with water; a precipitate of oxychloride occurs, the formation of which has been explained under the similar synthetical reaction. The occurrence of this precipitate distinguishes antimony from arsenicum, but is a reaction that cannot be fully relied upon in analysis, because requiring the presence of too much material and the observance of too many conditions. Add a sufficient quantity of hydrochloric acid to dissolve the precipitate, and boil a piece of copper in the solution as directed in the corresponding test for arsenicum (*vide* page 191); antimony is

deposited on the copper. Wash, dry, and heat the copper in a test-tube as before; the antimony, like the arsenicum, is volatilized off the copper and condenses on the side of the tube as white oxide; but the sublimate, from its low degree of volatility, condenses close to the copper; moreover, it is destitute of crystalline character,—that is to say, it is *amorphous* (*a, a*, without; *μορφή, morphê*, shape).

Shake out the copper and boil water in the tube for several minutes. Do the same with the arsenical sublimate similarly obtained. The deposit of arsenic slowly dissolves, and may be recognised in the solution by ammonio-nitrate of silver; the antimonial sublimate is insoluble.

Third Analytical Reaction.—Perform the experiments described under Marsh's test for arsenicum (pp. 192–3), carefully observing all the details there mentioned, but using a few drops of solution of chloride of antimony or tartar emetic instead of the arsenical solution. Antimoniuretted hydrogen, or hydride of antimony (SbH_3), is formed and decomposed in the same way as arseniuretted hydrogen.

To one of the arsenicum spots on the porcelain lid (p. 193) add a drop of solution of "chloride of lime" (bleaching-powder); it quickly dissolves. Do the same with an antimony spot; it is unaffected.

Heat more quickly causes the volatilization of an arsenicum than an antimony spot; sulphhydrate of ammonium more readily dissolves the antimony than the arsenicum.

Boil water for several minutes in the beaker or wide test-tube containing the arsenious sublimate (p. 193); it slowly dissolves and may be recognised in the solution by the yellow precipitate given on the addition of solution of ammonio-nitrate of silver. The antimonial sublimate, similarly treated, gives no corresponding reaction.

Pass a slow current of sulphuretted hydrogen through the delivery-tube removed from the hydrogen-apparatus (page 193), and, when the air may be considered to have been expelled from the tube, gently heat that portion containing

the deposit of arsenicum; the latter will be converted into a *yellow* sublimate of sulphide of arsenicum. Remove the tube from the sulphuretted-hydrogen apparatus, and repeat the experiment with a similar antimony deposit; it is converted into *orange* sulphide of antimony, which, moreover, owing to inferior volatility, condenses nearer to the flame than sulphide of arsenicum.

Pass dry hydrochloric acid gas through the two delivery-tubes. This is accomplished by adapting first one tube and then the other by a cork to a test-tube containing a few lumps of common salt, on which a little sulphuric acid is poured during the momentary removal of the cork. The sulphide of antimony dissolves and disappears; the sulphide of arsenicum is unaffected.

Thorough perception of the chemistry of arsenicum and antimony will be obtained on constructing equations or diagrams descriptive of each of the foregoing reactions.

Antidote.—The introduction of poisonous doses of antimonials into the stomach is fortunately quickly followed by vomiting. If vomiting has not occurred, or apparently to an insufficient extent, any form of tannic acid may be administered (infusion of tea, nutgalls, cinchona, oak-bark, or other astringent solutions or tinctures), an insoluble tannate of antimony being formed, and absorption of the poison consequently somewhat retarded. The stomach-pump must be applied as quickly as possible.

Recently precipitated moist ferric hydrate is also, according to T. and H. Smith, a perfect absorbent of antimony from its solutions, the chemical action being probably, they say, similar to that which takes place between ferric hydrate and arsenious anhydride. It may be given in the form of a mixture of perchloride of iron with either carbonate of sodium or other soluble carbonate or bicarbonate or magnesia.

These statements may be verified by mixing together the various substances, filtering, and testing the filtrate for antimony in the usual manner.

DIRECTIONS FOR APPLYING THE FOREGOING REACTIONS TO THE ANALYSIS OF AN AQUEOUS SOLUTION OF A SALT OF ONE OF THE ELEMENTS ARSENICUM AND ANTIMONY.

Acidify the liquid with hydrochloric acid, and pass through it sulphuretted hydrogen :—

A *yellow* precipitate indicates arsenicum ;

An *orange* precipitate indicates antimony.

The result may be confirmed by the application of other tests.

DIRECTIONS FOR APPLYING THE FOREGOING REACTIONS TO THE ANALYSIS OF AN AQUEOUS SOLUTION OF SALTS OF BOTH ARSENICUM AND ANTIMONY.

Acidify a small portion of the liquid with hydrochloric acid, and pass through it sulphuretted hydrogen.

Note I.—If the precipitate by sulphuretted hydrogen is unmistakably orange, antimony may be put down as present, and arsenicum only further sought by the application of Fleitmann's test to the solution of the sulphides in aqua regia * freed from sulphur by boiling, or, better, to the original solution.

Note II.—Sulphide of antimony is far less readily soluble than sulphide of arsenicum in solution of carbonate of ammonium. But this fact possesses limited analytical value ; for the colour of the sulphides is already sufficient to distinguish the one from the other when they are unmixed ; and when mixed, much sulphide of antimony will prevent a little sulphide of arsenicum from being dissolved by the alkaline carbonate, while much sulphide of arsenicum will carry a little sulphide of antimony into the solution. When the proportions are, apparently, from the colour of the precipitate, less wide, solution of carbonate of ammonium will be found useful in roughly separating the one sulphide from the other. On filtering and neutralizing the alkaline solution by an acid,

* *Aqua Regia* is a mixture of four parts hydrochloric and three parts nitric acid. It was so called from its property of dissolving gold, the "king" of metals. Diluted with rather more than four times its bulk of water, it forms the *Acidum Nitro-hydrochloricum Dilutum*, B.P.

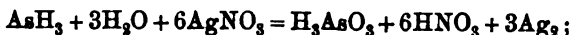
the yellow sulphide of arsenicum is reprecipitated. The orange sulphide of antimony will remain on the filter.

Note III.—Solution of bisulphite of potassium is said by Wöhler to be a good reagent for separating the sulphides of arsenicum and antimony, the former being soluble, the latter insoluble in the liquid.

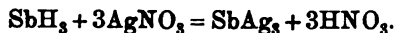
Note IV.—Another reagent for separating the sulphides of arsenicum and antimony is *strong* hydrochloric acid. As little water as possible must be present. On boiling, the sulphide of antimony dissolves, while the sulphide of arsenicum remains insoluble. The liquid slightly diluted, filtered, more water added, and sulphuretted hydrogen again transmitted, gives orange sulphide of antimony. The process should previously be tried on the precipitated mixed sulphides. The presence of arsenicum may be confirmed by the application of Fleitmann's test to the original solution.

Note V.—If the precipitate by sulphuretted hydrogen is unmistakably yellow, arsenicum may be put down as present, and any antimony detected by the previous or one of the following two processes. These two processes are rather long, and require much care in their performance; but are useful, because a small quantity of antimony in much arsenicum, or *vice versâ*, may be detected by their means.

First process.—Generate hydrogen and pass it through a small wash-bottle containing solution of acetate of lead, to free the gas from any trace of sulphuretted hydrogen it may possess, and then through a dilute solution of nitrate of silver contained in a test-tube. When the apparatus is in good working order, pour into the generating-bottle the solution to be examined, adding it gradually to prevent violent action. After the gas has been passing for five or ten minutes, examine the contents of the nitrate-of-silver tube; arsenicum, if present, will be found in the solution in the state of arsenious acid,



while antimony, if present, will be found in the black precipitate that has fallen, according to the following equation :



The arsenious radical may be detected in the clear, filtered, supernatant liquid, which still contains much nitrate of silver, by cautiously neutralizing with a very dilute solution of ammonia, or by adding a few drops of solution of ammonio-nitrate of silver, yellow arsenite of silver being produced. The antimony may be detected by washing the black precipitate, boiling it in an open dish with solution of tartaric acid, filtering, acidulating with hydrochloric acid, and passing sulphuretted hydrogen through the solution,—the orange sulphide of antimony being precipitated. (Hofmann.)

Second process.—Obtain the metallic deposit in the middle of the delivery-tube as already described under Marsh's test. Act on the deposit by sulphuretted hydrogen gas, and then by hydrochloric acid gas, as detailed in the third analytical reaction of antimony (p. 207). If both arsenicum and antimony are present, the deposit, after the action of sulphuretted hydrogen, will be found to be of two colours, the yellow sulphide of arsenicum being usually farther removed from the heated portion of the tube than the orange sulphide of antimony. Moreover, subsequent action of hydrochloric acid gas causes disappearance of the antimonial deposit, which is converted into chloride of antimony and carried off in the stream of gas.

The chief objection to this process is the liability of the operator mistaking sulphur, deposited from the sulphuretted hydrogen gas by heat, for sulphide of arsenicum. But the presence or absence of arsenicum is easily confirmed by applying Fleitmann's test to the original solution, while the process is most useful for the detection of a small quantity of a salt of antimony when mixed with much arsenical compounds.

The laboratory student may now proceed to the analysis of aqueous solutions of salts of any of the metallic elements

hitherto considered. The method followed may be that for the separation of the previous three groups, sulphuretted hydrogen being first passed through the solution to throw out arsenicum and antimony. The whole scheme of analysis is given on the next page. Three or four solutions should be examined before proceeding to the last group of metals.

Learners who have no opportunity of working at practical analysis will gain much knowledge by endeavouring, not to remember, but to understand these methods of separating elements from each other in a solution containing several compounds.

QUESTIONS AND EXERCISES.

267. What is the composition and source of the *Black Antimony* of pharmacy?

268. In what alloys is metallic antimony a characteristic ingredient?

269. What is the quantivalence of antimony as far as indicated by the formulæ of the official preparations?

270. By a diagram show how "*Butter of Antimony*" is prepared.

271. Write out equations or diagrams expressive of the reactions which occur in converting chloride of antimony into oxide.

272. What is the formula of *Tartar Emetic*?

273. Explain the official process for the preparation of *Oxy-sulphide of Antimony* (*Antimonium Sulphuratum*, B. P.) by aid of diagrams.

274. Give a comparative statement of the tests for arsenicum and antimony.

275. How is antimony detected in the presence of arsenicum?

276. How may arsenicum and iron be distinguished analytically?

277. Describe a method by which antimony, magnesium, and iron may be separated from each other.

278. Draw out an analytical chart for the examination of an aqueous liquid containing salts of arsenicum, zinc, calcium, and ammonium.

COPPER, MERCURY, LEAD, SILVER.

These metals, like arsenicum and antimony, are precipitated from acidified solutions by sulphuretted hydrogen, in the form

TABLE OF SHORT DIRECTIONS FOR THE ANALYSIS OF AN AQUEOUS SOLUTION OF SALTS OF ANY OR ALL OF THE METALLIC ELEMENTS HITHERTO CONSIDERED.
Acidity with HCl, and pass H₂S through the solution; filter.

Precipitate As Sb. Wash, separate by carbonate of ammonium, or by strong hydrochloric acid; or dissolve in a few drops of hydrochloric and nitric acids, and examine by Hofmann's method (p. 210).	Filtrate Fe Al Zn Ba Ca Mg K Na Am. Add AmHO*; AmHS; stir, filter.						
	Precipitate† Fe Al Zn. Wash, dissolve in HCl, boil with HNO ₃ (p. 183), add KHO in excess †, stir, filter.		Filtrate Ba Ca Mg K Na Am. Add Am ₂ CO ₃ , boil, filter.				
	Precipitate Fe. Examine orig. solution for ferrous or ferric state.	Filtrate Al Zn. Neut. by HCl, add AmHO, stir, filter.		Precipitate Ba Ca.		Precipitate Mg.	Filtrate Am Na K (page 136).
		Ppt. Al.	Sol. Zn.				

* It is not usually necessary here to add NH₄Cl to keep Mg in solution, the HCl with some of the NH₄HO commonly giving sufficient NH₄Cl for this purpose.

† Much time may sometimes be saved by carefully remembering the colour of the various hydrates and sulphides precipitated. Thus, if the AmHS precipitate is white, iron cannot be present, and AmHO, for Al and Zn, may be at once added to the hydrochloric solution of the precipitate. The *group-tests* in this Table are H₂S, AmHS, and Am₂CO₃.

‡ The pupil should vary the process sometimes, as described in the footnotes on pages 180 and 188. The treatment of other groups should occasionally be varied. By remembering the reactions on which analytical processes are based, a pupil may often devise modified processes, and thus frequently save time, avoid working in one groove only, and acquire broader chemical views.

of sulphides; but the sulphides, unlike those of arsenicum and antimony, are insoluble in alkalis. The atom of copper is usually bivalent, Cu'' ; mercury bivalent in the mercuric salts, Hg'' , and univalent in the mercurous salts, Hg' ; lead sometimes quadrivalent, Pb'''' , but generally exerting only bivalent activity, Pb'' ; and silver univalent, Ag' .

COPPER.

Symbol Cu. Atomic weight 63·5.

Source.—The commonest ore of this metal is *copper pyrites*, a double sulphide of copper and iron, raised in Cornwall; Australia and Russia supply *malachite*, a mixed carbonate and hydrate; much ore is also imported from South America. It is smelted in enormous quantities at Swansea, South Wales, a locality peculiarly fitted for the operation on account of its proximity to the coal-fields, and its position as a sea-coast town—these advantages insuring cheap fuel and freightage to the different metallurgical establishments. An economical method of smelting copper pyrites and other sulphides has recently been introduced by Hollway. After the sulphide is once melted air is driven, not over, as usual, but through, the mass; the combustion of the sulphur then becomes self-supporting, and is greatly accelerated.

Alchemy.—The alchemists termed this metal *Venus*, perhaps on account of the beauty of its lustre, and gave it her symbol ☿, a compound hieroglyphic also indicating a mixture of gold ☉ and a certain hypothetical substance called *acrimony* ☿, the corrosive nature of which was symbolized by the points of a Maltese cross. To this day the blue show-bottle in the shop-window of the pharmacist is occasionally ornamented by such a symbol, indicative, possibly, of the fact that the blue liquid in the vessel is a preparation of copper.

Coinage.—The material of our "copper" coinage is now a bronze mixture composed, in 100 parts by weight, of 95 copper, 4 tin, and 1 zinc, the same as in the copper coinage of France. The penny is coined at the rate of 48 pence in one pound avoirdupois of 7000 grains, or 453·6 grammes; the halfpenny at 80 in the pound avoirdupois, and the farthing at 160. British bronze coins are a legal tender in payments to the amount of 1s.

Metallic Copper (*Cuprum*, B. P.) in the form of fine wire, about No. 25, is used in preparing *Spiritus Ætheris Nitrosi*,

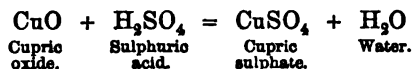
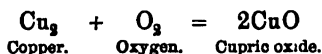
B. P. *Copper Foil*, B. P., is "pure metallic copper, thin and bright."

Quantivalence.—Copper forms two classes of salts; in one the atom is bivalent (Cu''), in the other it exerts univalent activity (Cu'). The former are of primary importance, the latter being for the most part unstable and wanting in technical interest. Their compounds are distinguished as cupric and cuprous; but those of the higher class only have general interest, and will be almost exclusively alluded to in the following paragraphs. Cuprous iodide (Cu_2I_2) will be subsequently referred to as a convenient form in which to remove iodine from solution, while the formation of cuprous oxide (Cu_2O), under given circumstances, will come under notice as an indicator of the presence of sugar in a liquid.

REACTIONS HAVING (a) SYNTHETICAL AND (b) ANALYTICAL INTEREST.

(a) *Synthetical Reactions.*

The processes for the following salts include the only synthetical reactions having any medical or pharmaceutical interest:—1, cupric oxide, the black oxide of copper, prepared by heating fragments of copper to low redness on a piece of earthenware in an open fire; 2, cupric sulphate, the common sulphate of copper, prepared by boiling black oxide and about an equal weight of sulphuric acid in water, filtering, and setting aside the solution so that crystals may form on cooling; and 3, ammonio-sulphates of copper, for the preparation of which see pages 198 and 217.



Sulphate of Copper (*Cupri Sulphas*, B. P.) ($\text{CuSO}_4, 5\text{H}_2\text{O}$), *blue vitriol*, *bluestone*, or *cupric sulphate*, is the only copper salt of much importance in Pharmacy. It is a by-product in silver-refining ($2\text{Ag}_2\text{SO}_4 + \text{Cu}_2 = 2\text{CuSO}_4 + 2\text{Ag}_2$). A little is formed in roasting copper pyrites. In the latter case, some sulphide of iron and sulphide of copper are oxidized to sulphates; but the low red heat finally employed decomposes

the sulphate of iron, while the sulphate of copper is unaffected; it is purified by crystallization from a hot aqueous solution, though frequently much sulphate of iron remains in the crystals. Sulphate of copper results on dissolving in diluted sulphuric acid the black oxide (CuO) obtained in annealing copper plates (*see* the foregoing equation); it may also be prepared by boiling copper with three times its weight of sulphuric acid ($2\text{H}_2\text{SO}_4 + \text{Cu} = \text{CuSO}_4 + \text{SO}_2 + 2\text{H}_2\text{O}$), diluting, filtering, evaporating, and crystallizing. In this process a little black sulphide of copper is formed.

Anhydrous Sulphate of Copper, B. P. (CuSO_4), is a yellowish white powder prepared by depriving the ordinary blue crystals of sulphate of copper of their water of crystallization by exposing to a temperature of about 400°F . It is used in testing alcohol and similar spirituous liquids for water, becoming blue if the latter be present.

Verdigris (from *verde-gris*, Sp., green-grey) is a Subacetate or Oxyacetate of Copper (B. P.) ($\text{Cu}_2\text{O}_2\text{C}_2\text{H}_3\text{O}_2$), obtained by exposing alternate layers of copper and fermenting refuse grape-husks to the action of air. Digested with twice its weight of acetic acid and a little water, the mixture being evaporated to dryness and the residue dissolved in water, it forms the official Solution of Acetate of Copper ($\text{Cu}_2\text{C}_2\text{H}_3\text{O}_2$).

The modes of forming *Cupric Sulphide*, *Hydrate*, *Oxide*, *Ferro-cyanide* and *Arsenite*, as well as *Metallic Copper*, are incidentally alluded to in the following analytical paragraphs.

(b) *Reactions having Analytical Interest (Tests).*

First Analytical Reaction.—Pass sulphuretted hydrogen through an acidified solution of a copper salt (sulphate, for example); black cupric sulphide (CuS) falls.

Second Analytical Reaction.—To an aqueous copper solution add sulphhydrate of ammonium; by this reagent, also, cupric sulphide is precipitated, insoluble in excess.

Note.—Cupric sulphide is not altogether insoluble in sulphhydrate of ammonium if free ammonia or much ammoniacal salt be present; it is quite insoluble in the fixed alkaline sulphhydrates.

Third Analytical Reaction.—Immerse a piece of iron or steel, such as the point of a penknife or a piece of wire, in a few

drops of copper solution ; the copper is deposited, of characteristic colour, an equivalent quantity of iron passing into solution.

By this reaction copper may be recovered on the large scale from waste solutions, old hoop or other scrap iron being thrown into the liquors.

Fourth Analytical Reaction.—Add ammonia to a cupric solution ; cupric hydrate (Cu_2HO) of a light-blue colour is precipitated. Add excess of ammonia ; the precipitate is redissolved, forming a blue solution of ammonio-salt of copper, so deep in colour as to render ammonia an exceedingly delicate test for this metal.

An ammonio-sulphate of copper may be obtained in large crystals by adding strongest solution of ammonia to powdered sulphate of copper until the salt is dissolved, placing the liquid in a test-glass or cylinder, cautiously pouring in twice its volume of strong alcohol or methylated spirit, taking care that the liquids do not become mixed, tying over the vessel with bladder, and setting aside for some weeks in a cool place. (Wittstein.) The constitution of ammonio-sulphate and other ammonio-salts of copper and corresponding salts of silver will be alluded to in connection with "white precipitate," the official "ammoniated mercury."

Fifth Analytical Reaction.—Add solution of potash or soda to a cupric solution ; cupric hydrate (Cu_2HO) is precipitated, insoluble in excess. Boil the mixture in the test-tube ; the hydrate is decomposed, losing the elements of water, and becoming the black anhydrous oxide (CuO).

Sixth Analytical Reaction.—Add solution of ferrocyanide of potassium (K_4Fcy) to an aqueous cupric solution ; a reddish-brown precipitate of cupric ferrocyanide (Cu_3Fcy) falls. This is an extremely delicate test for copper.

Seventh Analytical Reaction.—To a cupric solution add solution of arsenic, and cautiously neutralize with alkali ; green cupric arsenite (CuHAsO_3) falls.

Note.—This precipitate has been already mentioned under arsenicum. An arsenicum salt is thus a test for copper, as a

copper salt is for arsenicum,—a remark that may obviously be extended to most analytical reactions; for the *body acted upon characteristically by a reagent is as good a test for the reagent as the reagent is for it*; indeed it becomes a reagent when the other body is the object of search.

Antidotes.—In cases of poisoning by compounds of copper, iron filings should be administered, the action of which has just been explained (*see* third analytical reaction). Ferrocyanide of potassium may also be given (*see* sixth analytical reaction). Albumen forms with copper a compound insoluble in water; hence raw eggs should be swallowed, vomiting being induced or the stomach-pump applied as speedily as possible.

QUESTIONS AND EXERCISES.

279. What are the analytical relations of copper, mercury, lead, and silver to each other and to arsenicum and antimony?

280. Name the sources of copper.

281. What proportion of copper is contained in English and French "copper" coins?

282. Give diagrams showing how Sulphate of Copper is prepared on the small and large scales.

283. Work out a sum showing how much Crystallized Sulphate of Copper may be obtained from 100 parts of sulphide.—*Ans.* 261½.

284. How may Oxide of Copper be prepared?

285. Mention the formula of Verdigris?

286. Name a good clinical test for copper.

287. What is the analytical position of copper?

288. Mention the chief tests for copper.

289. How may copper be separated from arsenicum?

290. Why is finely divided iron an effective antidote in cases of poisoning by copper?

MERCURY.

Symbol Hg. Atomic weight 200.

Molecular weight 200 (*not* double the atomic weight).

Source.—Mercury occurs in nature as sulphide (HgS), forming the ore *cinnabar* (an Indian name expressive of some-

thing red), and is obtained from Spain, California, Eastern Hungary, China, Japan, and Peru.

Preparation.—The metal is separated by roasting off the sulphur and then distilling, or, better, distilling with lime, which combines with and retains the sulphur.

Properties.—Mercury (*Hydrargyrum*, B.P.) is a silver-white, lustrous metal, liquid at common temperatures. It boils at 662° F., and at -40° F. solidifies to a malleable mass of octahedral crystals. When quite free from other metals it does not tarnish, and its globules roll freely over a sheet of white paper without leaving any streak or losing their spherical form.

Formula.—The formula of the mercury molecule is Hg and not Hg₂, because (at all events at the high temperature at which alone the weight of its vapour can be determined), two volumes, which if hydrogen would weigh two parts (H₂) or oxygen thirty-two parts (O₂), in the case of mercury vapour weigh only two hundred parts (Hg); that is, only once the atomic weight, not twice. That 200, and not 100, is the atomic weight of mercury is shown by the fact that 200 is the minimum proportion, relative to 1 of hydrogen, in which mercury combines, and by its relations to heat. Still it is difficult to imagine an atom existing in the free state in nature; and the suggestion has been made that (as is proved to be the case with sulphur) mercury, as we know it, is in abnormal condition, and that if the weight of its vapour could be taken at a lower temperature or under some other condition, its molecular weight might be found to be 400. Similar remarks may be made respecting zinc, the molecular weight of which, so far as we know, is identical with its atomic weight.

Medicinal Compounds.—The compounds of mercury used in medicine are all obtained from the metal. The metal itself, rubbed with chalk or with confection of roses and powdered liquorice-root, or with lard and suet, until globules are not visible to the unaided eye, is often used in medicine. The preparations are:—the *Hydrargyrum cum Oreta*, B. P., or "Grey Powder;" *Pilula Hydrargyri*, B. P., or "Blue Pill;" and *Unguentum Hydrargyri*, B. P., or "Blue Ointment." There are also a Compound Ointment, a Plaster of Mercury, a Plaster of Ammoniacum and Mercury, a Liniment, and a Suppository, all official. Their therapeutic effects are probably due, not to the large quantity of metallic mercury in them, but, to the small quantities of black and red oxide which

occur in them through the action of the oxygen of the air on the finely divided metal. The proportion of oxide or oxides varies according to the age of the specimen.

All these medicinal preparations of metallic mercury are indefinite and unsatisfactory, and that through no fault of the pharmacist. They much need investigation by therapeutists. Here, as in many similar cases, if medicine would first ascertain her own requirements, and then make them known, her handmaid pharmacy would be found to be quite capable of supplying them.

Mercurous and Mercuric Compounds.—Mercury combines with other elements and radicals in two proportions: those compounds in which the other, acidulous, radicals are in the lesser amount are termed *mercurous*, the higher being *mercuric*. Thus calomel (HgCl)* is mercurous chloride, while corrosive sublimate (HgCl_2) is mercuric chloride. In every pair of mercury compounds the mercuric contains *twice* as much complementary radical, in proportion to the mercury, as the mercurous.

Note on Nomenclature.—The remarks made concerning the two classes of iron salts, ferrous and ferric (p. 155), apply in the main to the two series of mercury salts. The latter are systematically distinguished in most modern works by the terms *mercurous* and *mercuric*. In the British Pharmacopœia, however, which includes only a few in comparison with the whole number of mercury salts, older and more strongly contrasted names are employed, thus:—

Systematic names.	Official names.
Mercurous iodide . . .	Green iodide of mercury.
Mercuric iodide . . .	Red iodide of mercury.
Mercurous nitrate . . .	Not mentioned in B. P.
Mercuric nitrate . . .	Nitrate of mercury.
Mercurous sulphate . . .	Not mentioned in B. P.
Mercuric sulphate . . .	Sulphate of mercury.
Mercurous chloride . . .	Subchloride of mercury.
Mercuric chloride . . .	Perchloride of mercury.
Mercurous oxide . . .	Black oxide of mercury.
Mercuric oxide . . .	Red oxide of mercury.

* The specific gravity of the vapour of calomel, and the fact that the salt is *not* decomposed at the temperature at which its specific gravity is taken, indicate that the formula of calomel is HgCl , and not Hg_2Cl_2 .

Specific Gravity.—Mercury is 13.6 times as heavy as water.

Amalgams.—The compound formed on fusing metals together is usually termed an *alloy* (*ad* and *ligo*, to bind); but if mercury is a constituent, an *amalgam* (μάλαγμα, *malagma*, from μαλάσσω, *malassō* to soften, the presence of mercury lowering the melting-point of such a mixture). Most metals, even hydrogen, according to Loew, form amalgams.

REACTIONS HAVING (a) SYNTHETICAL AND (b) ANALYTICAL INTEREST.

(a) *Synthetical Reactions.*

The two Iodides.

First Synthetical Reaction.—Rub together a small quantity of mercury and iodine, controlling the rapidity of combination by adding a few drops of spirit of wine, which, by evaporation, absorbs heat, and thus keeps down temperature. The product is either mercuric iodide, mercurous iodide, or a mixture of the two, as well as mercury or iodine if excess of either has been employed. If the two elements have been previously weighed in single atomic proportions, 200 of mercury to 127 of iodine (about 8 to 5, or 1 ounce of mercury to 278 grains of iodine), the mercurous or green (greyish green) iodide results, HgI (*Hydrargyri Iodidum Viride*, B. P.); if in the proportion of one atom of mercury to two atoms of iodine (200 to twice 127, or about 4 to 5), the mercuric or red iodide, HgI_2 , results, an iodide that is also official, but made in another way (*see* p. 223). The green iodide should be made and dried (without heat) with as little exposure to light as possible.

Mercurous iodide is decomposed slowly by light, and quickly by heat, into mercuric iodide and mercury. Mercuric iodide occurring as an impurity in mercurous iodide may be detected by digesting in ether (in which mercurous iodide is insoluble), filtering, and evaporating to dryness; mercuric iodide remains. Mercuric iodide is stable, and may be sublimed in scarlet crystals without decomposition. (For details

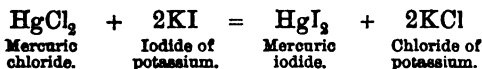
of the method by which a specimen of the crystals may be obtained, and the precautions to be observed, *vide* "corrosive sublimate," p. 226).

Relation of Mercuric Iodide to Light.—In condensing, mercuric iodide is at first yellow, afterwards acquiring its characteristic scarlet colour. This may be shown by smearing or rubbing a sheet of white paper with the red iodide, and then holding the sheet before a fire or over a flame for a few seconds. As soon as the paper becomes hot the red instantly changes to yellow, and the salt does not quickly regain its red colour, even when cold, if the paper is carefully handled. But if a mark be made across the sheet by anything at hand, or the salt be pressed or rubbed in any way, the portions touched immediately return to the scarlet condition. According to Warrington, this change is consequent upon rhomboidal crystals being converted into octahedra with a square base, and will serve as an excellent illustration of the influence of physical structure in causing colour. The yellow modification so acts on the rays of white light shining on its particles as to absorb the violet and reflect the complementary hue, the yellow, which, entering the eye of the observer, strikes his retina, and thus conveys to the brain the impression of yellowness; and the red modification, though actually the same chemical substance, is sufficiently different in the structure of its particles to absorb the green constituent of white light and reflect the complementary ray, the red.

Illustration of the Chemical law of Multiple Proportions (p. 45).—Applying the atomic theory to the above iodides, it will at once be apparent why mercury and iodine should combine in the proportion of 200 of mercury with either 127 or 254 of iodine, and not with any intermediate quantity. For it is part of that theory that masses are composed of atoms, and that atoms are indivisible, and that the weight of the atom of mercury is to that of iodine as 200 is to 127. Mercury and iodine can only combine, therefore, in atomic proportions, atom to atom (which is the same as 200 to 127), or one atom to two atoms (which is the same as 200 to 254). To attempt to combine them in any intermediate proportion would be useless, a mere mixture of the two iodides would result. A higher proportion of mercury than 200 to 127 of iodine gives but a mixture of mercurous iodide and mercury; a higher proportion of iodine than 254 to 200 of mercury gives but a mixture of mercuric iodide and iodine. Or, for example, 200

grains of mercury mixed with, say, 200 of iodine would yield 139 grains of mercurous iodide and 261 grains of mercuric iodide; for the 200 grains of mercury uniting with 127 grains of the iodine gives, for the moment, 327 grains of mercurous iodide and 73 grains of iodine still free. The 73 grains of iodine will immediately unite with 188 grains of the mercurous iodide (for if 127 of I require 327 of HgI to form HgI₂, 73 will require 188), and form 261 grains of mercuric iodide, diminishing the 327 grains of mercurous iodide to 139 grains.

Preparation of Red Iodide of Mercury by precipitation.—To a few drops of a solution of a mercuric salt (corrosive sublimate, for example) add solution of iodide of potassium, drop by drop; a precipitate of mercuric iodide, HgI₂, forms, and at first quickly redissolves, but is permanent when sufficient iodide of potassium has been added. Continue the addition of iodide of potassium; the precipitate is once more redissolved.

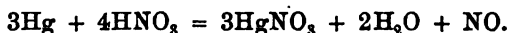


Notes.—When first precipitated, mercuric iodide is yellowish red, but soon changes to a beautiful scarlet. Its solubility either in solution of the mercuric salt or in solution of iodide of potassium renders the detection of a small quantity of a mercuric salt by iodide of potassium, or a small quantity of an iodide by a mercuric solution, difficult, and hence lessens the value of the reaction as a test. But the reaction has synthetical interest, the method by precipitation being that adopted in the British Pharmacopœia (*Hydrargyri Iodidum Rubrum*, B. P.). Mercuric iodide thus made has the same composition as that prepared by direct combination of its elements. Equivalent proportions of the two salts must be used in making the preparation (HgCl₂ = 271; 2KI = 332). About 4 parts of corrosive sublimate are dissolved in 50 or 60 of water (warmth quickens solution) and 5 of iodide of potassium in 15 or 20 of water, the solutions mixed and the precipitate collected on a filter, drained, washed twice with distilled water, and dried on a plate over a water-bath. Mercuric nitrate, which is more soluble and therefore some-

what more convenient for use on the large scale, may be used instead of the mercuric chloride. The mercury in mercuric or mercurous iodide is set free and sublimes in globules on heating either powder with dried carbonate of sodium in a test-tube; the iodine may be detected by digesting with solution of soda, filtering, and to the solution of iodide of sodium thus formed adding starch paste and acidulating with nitrous acid, when blue iodide of starch results. Mercuric iodide is insoluble in water, slightly soluble in alcohol, tolerably soluble in ether. Precipitated red iodide of mercury mixed with white wax, lard, and oil forms the *Unguentum Hydrargyri Iodidi Rubri*, B. P. *Donovan's Solution* contained mercuric and arsenious iodides.

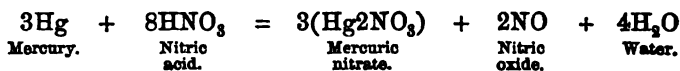
The two Nitrates.

Second Synthetical Reaction.—Mix a little nitric acid in a test-tube with four or five times its bulk of water, add a small globule of mercury, and set the tube aside for a few hours, in a cool place; solution of mercurous nitrate (HgNO_3) will be formed, and nitric oxide (NO) evolved. The solution may be retained for subsequent analytical operations.



Third Synthetical Reaction.—Place mercury in strong nitric acid, and warm the mixture; mercuric nitrate is formed, and will be deposited in crystals as the solution cools. Retain the product for a subsequent experiment.

The mercuric nitrates vary somewhat in composition, according to the proportion, strength, and temperature of the acid used in their formation. A mercuric nitrate may be obtained having the formula Hg_2NO_3 .



Mercuric oxynitrates.—From the normal mercuric nitrate several oxynitrates may be obtained. Thus on merely evaporating a solution of mercuric nitrate, and cooling, crystals having the formula $\text{Hg}_3\text{O}_3\text{NO}_3$ are deposited. The latter, by

washing with cold water, yield a *yellow* pulverulent oxynitrate, $\text{Hg}_6\text{O}_4\text{4NO}_3$: mixed with lard, this has sometimes been used as an ointment. Boiled in water, the yellow gives a *brick-red* oxynitrate, $\text{Hg}_3\text{O}_5\text{2NO}_3$.

The Pharmacopœial preparations of mercuric nitrate are *Liquor Hydrargyri Nitratus Acidus* and *Unguentum Hydrargyri Nitratis*. The former is made by placing four ounces of mercury in five fluid ounces of nitric acid diluted with an ounce and a half of water, and, when the metal is dissolved, boiling gently for fifteen minutes.

The two Sulphates.

Fourth Synthetical Reaction.—Boil two or three grains of mercury with a few drops of strong sulphuric acid in a test-tube, or, better, small dish; sulphurous acid gas (SO_2) is evolved, and mercuric sulphate (*Hydrargyri Sulphas*, B. P.) (HgSO_4) results—a white heavy crystalline powder.



Between two and three ounces of mercuric sulphate may be prepared from a fluid drachm of mercury and a fluid ounce of sulphuric acid boiled together in a small dish. These are the official proportions. The operation is completed and any excess of acid removed by evaporating the mixture of metal and liquid to dryness, either in the open air or in a fume-chamber, sulphuric acid vapours being excessively irritating to the mucous membrane of the nose and throat; dry crystalline mercuric sulphate remains. If residual particles of mercury are observed, the mass should be damped with sulphuric acid and again heated.

By-products.—In chemical manufactories, secondary products, such as the sulphurous gas of the above reaction, are termed *by-products*, and, if of value, are utilized. In the present case the gas is of no immediate use, and is therefore allowed to escape. When very pure sulphurous acid gas is required for experiments on the small scale, this would be the best method of making it, a delivery-tube being adapted by a

cork to the mouth of a flask containing the acid and metal. The sulphate of mercury would then become the by-product.

Mercuric oxysulphate.—Water decomposes mercuric sulphate into a soluble acid salt and an insoluble yellow oxysulphate ($\text{Hg}_3\text{O}_3\text{SO}_4$). The latter is called *Turpeth mineral*, from its resemblance in colour to the powdered root of *Ipomœa turpethum*, an Indian substitute for jalap. The yellow sulphate of mercury was formerly official, but is now seldom used.

Fifth Synthetical Reaction.—Rub a portion of the dry mercuric sulphate of the previous reaction with as much mercury as it already contains; the product, when the two have thoroughly blended, is mercurous sulphate (Hg_2SO_4): it may be retained for a subsequent experiment.

Molecular Weight.—The exact proportion of mercury to sulphate is merely a matter of calculation; for the combining proportion of a compound (if it possess any combining-power) is the sum of the combining proportions of its constituents. *In other words, the combining weight of a molecule is simply the sum of the weights of its constituent atoms*, or, more generally, *the molecular weight of a compound is the sum of the atomic weights of its elements*. In accordance with this rule (sometimes called the fourth law of chemical combination, though only a deduction from the first—p. 43), 296 of mercuric sulphate and 200 of mercury (about 3 to 2) are the exact proportions necessary to the formation of mercurous sulphate.

The two Chlorides.

Sixth Synthetical Reaction.—Mix thoroughly a few grains of dry mercuric sulphate with about four-fifths its weight of chloride of sodium, and heat the mixture slowly in a test-tube in a fume-chamber or in the open air to leeward of the operator; mercuric chloride (HgCl_2), or *corrosive sublimate* (*Hydrargyri Perchloridum*, B. P.), sublimes and condenses in the upper part of the tube in heavy colourless crystals or a crystalline mass.

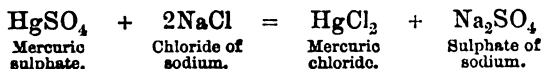
Somewhat larger quantities (in the proportion of 20 of sulphate to 16 of salt and, *vide infra*, 1 of black oxide of man-

ganese) may be sublimed in a pair of two-ounce or three-ounce round-bottom gallipots, the one inverted over the other, and the joint luted by moist fireclay (the powdered clay kneaded with water to the consistence of dough). The luting having been allowed to dry (somewhat slowly to avoid cracks), the pots are placed upright on a sand-tray (plate-shape answers very well), sand piled round the lower and a portion of the upper pot, and the whole heated over a good-sized gas-flame (if an air gas-flame it should be an inch and a half wide and four or five inches long) for an hour or more (see fig. 36—pots raised to show joint). Red Iodide of Mercury and Calomel may be sublimed in the same way. The former requires less, the latter more, heat than corrosive sublimate.

Fig. 36.



SUBLIMATION.



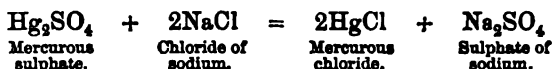
Note.—If the mercuric sulphate contain any mercurous sulphate, some calomel may be formed. This result will be avoided if 2 or 3 per cent. of black oxide of manganese be previously mixed with the ingredients, the action of which is to eliminate chlorine from the excess of chloride of sodium used in the process, the chlorine converting any calomel into corrosive sublimate. Manganate of sodium and a lower oxide of manganese are simultaneously produced.

Precaution.—The operation is directed to be conducted with care in a fume-chamber or in the open air, because the vapour of corrosive sublimate, which might possibly escape, is very acrid and highly poisonous. Its vulgar name is indicative of its properties.

Ten grains of perchloride of mercury and the same quantity of chloride of ammonium in one pint of water form the *Liquor Hydrargyri Perchloridi*, B. P. A dilute aqueous solution of perchloride of mercury is liable to decomposition, calomel

being precipitated, water decomposed, hydrochloric acid formed, and oxygen gas evolved. The presence of excess of chloride of ammonium, with a portion of which the mercuric chloride forms a stable double salt, prevents the decomposition.

Seventh Synthetical Reaction.—Mix a few grains of the mercurous sulphate of the fifth reaction with about a third of its weight of chloride of sodium, and sublime in a test-tube; crystalline mercurous chloride (HgCl) or calomel (*Hydrargyri Subchloridum*, B. P.) results. Larger quantities may be prepared in the manner directed for corrosive sublimate, a somewhat higher temperature being employed: similar precautions must also be observed. The official proportions are 10 of mercuric sulphate to 7 of mercury and 5 of dry chloride of sodium. "Moisten the sulphate of mercury with some of the water, and rub it and the mercury together until globules are no longer visible; add the chloride of sodium, and thoroughly mix the whole by continued trituration. When dry, sublime by a suitable apparatus into a chamber of such size that the calomel, instead of adhering to its sides as a crystalline crust, shall fall as a fine (dull-white) powder on its floor. Wash this powder with boiling distilled water until the washings cease to be darkened by a drop of sulphhydrate of ammonium. Finally, dry at a heat not exceeding 212°F. , and preserve in a jar or bottle impervious to light."



The term *calomel* (*καλός, kalos*, good, and *μέλας, melas*, black) is said to relate to the use of the salt as a *good* remedy for *black bile*, but probably was simply indicative of the esteem in which black sulphide of mercury was held, the compound to which the name calomel was first applied.

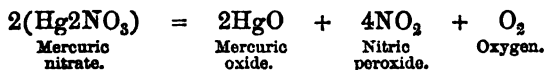
Test for corrosive sublimate in Calomel.—If the mercurous sulphate contains mercuric sulphate, some mercuric chloride will also be formed. Corrosive sublimate is soluble in water, calomel insoluble; the presence of the former may therefore be proved by boiling a few grains of the calomel in distilled

water, filtering and testing by sulphuretted hydrogen or sulphhydrate of ammonium as described hereafter. If corrosive sublimate is present, the whole bulk of the calomel must be washed with hot distilled water till the filtrate ceases to give any indications of mercury. Corrosive sublimate is more soluble in alcohol, and still more in ether, calomel insoluble. Ether in which calomel has been digested should therefore, after filtration, yield no residue on evaporation. Calomel is converted by hydrocyanic acid into mercuric salt, and a black powder readily yielding metallic mercury. Powell and Bayne have shown that a certain proportion of hydrochloric acid arrests this action.

Note.—The above process is that of the British Pharmacopœia; but calomel may also be made by other methods. Calomel mixed with lard forms the *Unguentum Hydrargyri Subchloridi*, B. P., and with sulphurated antimony, guaiacum resin, and castor-oil, the *Pilula Hydrargyri Subchloridi Composita*, B. P., or “Plummer’s Pill.”

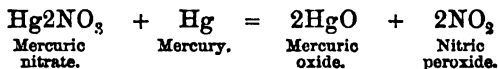
The two Oxides.

Eighth Synthetical Reaction.—Evaporate the mercuric nitrate of the third reaction to dryness in a small dish, in a fume-chamber or in the open air if more than a few grains have been prepared, and heat the residue till no more fumes are evolved; mercuric oxide (HgO), “Red Precipitate,” the Red Oxide of Mercury (*Hydrargyri Oxidum Rubrum*, B. P.), remains.



The nitric constituents of the salt may be partially economized by previously thoroughly mixing with the dry mercuric nitrate as much mercury as is used in its preparation, or as much as it already contains (ascertained by calculation from the atomic weights and the weight of nitrate under operation, as in making mercurous sulphate), and well heating the mixture. In this case the free mercury is also converted into mercuric oxide. This is the official process, the Pharmacopœial quantities being four ounces of mercury dissolved in four and a half fluid ounces of nitric acid diluted with two ounces of water, the solution evaporated to dryness, the

residue thoroughly mixed with four more ounces of mercury, and the whole heated until acid vapours cease to be evolved. (Mercuric oxide is tested for nitrate by heating a little of the sample in a test-tube, when orange nitrous vapours are produced and are visible in the upper part of the tube, if nitrate be present.)



Mercuric oxide is an orange-red powder, more or less crystalline according to the extent to which it may have been stirred during preparation from the nitrate, much rubbing giving the crystals a pulverulent character. Mixed with yellow wax and oil of almonds it yields the *Unguentum Hydrargyri Oxidi Rubri*, B. P. (1 part in 8). Mercuric oxide, in contact with oxidizable organic matter, is liable to reduction to black or mercurous oxide.

Ninth Synthetical Reaction.—To solution of potash, or soda, or lime-water, in a test-tube or larger vessel, add solution of corrosive sublimate or of mercuric nitrate; yellow oxide of mercury, or mercuric oxide (HgO), is precipitated (*Hydrargyri Oxidum Flavum*, B. P.).

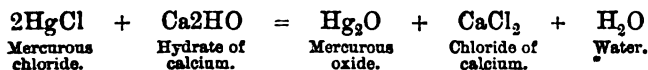


Eighteen grains of corrosive sublimate to ten ounces of lime-water form the *Lotio Hydrargyri Flava*, B. P. The precipitate only differs physically from the red mercuric oxide; the yellow is in a more minute state of division than the red. Mercuric oxide is very slightly soluble in water, but sufficiently so to communicate a decidedly metallic taste.

Tenth Synthetical Reaction.—To calomel add solution of potash or soda, or lime-water; black oxide of mercury, or mercurous oxide (Hg_2O), is produced, and may be filtered off, washed, and dried. (This reaction and the formation of a white curdy precipitate, on the addition of solution of nitrate of silver to the filtrate from the mercurous oxide, acidified by

nitric acid, form sufficient evidence of a powder being or containing calomel. The curdy precipitate is chloride of silver.)

Thirty grains of calomel to ten ounces of lime-water form the *Lotio Hydrargyri Nigra*, B. P.



(b) *Analytical Reactions (Tests).*

(The mercury occurring as mercuric or mercurous salt.)

First Analytical Reaction.—The Copper Test. Deposition of mercury upon, and sublimation from copper.—Place a small piece of bright copper, about half an inch long and a quarter of an inch broad, in a solution of any salt of mercury, mercurous or mercuric, and heat in a test-tube; the copper becomes coated with mercury in a fine state of division. (The absence of any notable quantity of nitric acid must be insured, or the copper itself will be dissolved. See below.) Pour away the supernatant liquid from the copper, wash the latter once or twice by pouring water into, and then out of, the tube, remove the metal, take off excess of water by gentle pressure in a piece of filter-paper, dry the copper by passing it quickly through a flame, holding it by the fingers; finally, place the copper in a dry, narrow test-tube, and heat to redness in a flame, the tube being held almost horizontally; the mercury sublimes and condenses as a whitish sublimate of minute globules on the cool part of the tube outside the flame. The globules aggregate on gently pressing with a glass rod, and are especially visible where flattened between the rod and the side of the test-tube.

Notes on the test.—This is a valuable test for several reasons:—It is very delicate when performed with care. It brings before the observer the element itself—one which, from its metallic lustre and fluidity, cannot be mistaken for any other. It separates the element both from mercurous and mercuric salts. Mercury can in this way readily be

eliminated in the presence of most other substances, organic or inorganic.

In performing the test the presence of any quantity of nitric acid may be avoided by adding an alkali until a slight permanent precipitate appears, and then reacidifying with a few drops of acetic or hydrochloric acid ; or by concentrating in an evaporating-dish after adding a little sulphuric acid, and then rediluting.

Tests, continued. (The mercury occurring as mercuric salt.)

Second Analytical Reaction.—To a few drops of a solution of a mercuric salt (corrosive sublimate, for example) add solution of iodide of potassium, drop by drop ; a precipitate of mercuric iodide, HgI_2 , forms, and at first quickly redissolves, but is permanent when sufficient iodide of potassium has been added. Continue the addition of iodide of potassium ; the precipitate is once more redissolved.

Note.—When first precipitated, mercuric iodide is yellowish red, but soon changes to a beautiful scarlet. Its solubility either in solution of the mercuric salt or in solution of iodide of potassium renders the detection of a small quantity of a mercuric salt by iodide of potassium, or a small quantity of an iodide by mercuric solution, difficult, and hence lessens the value of the reaction as a test.

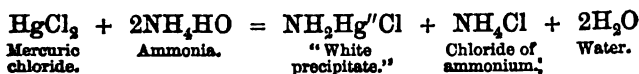
Third Analytical Reaction.—Add a solution of mercuric salt to solution of ammonia, taking care that the mixture, after well stirring, still smells of ammonia ; a white precipitate falls.

Ammoniated Mercury.

Performed in a test-tube, this reaction is a very delicate test of the presence of a mercuric salt ; performed in larger vessels, the mercuric salt being corrosive sublimate (3 ounces dissolved in 3 pints of distilled water, the solution poured into 4 fluid ounces of Solution of Ammonia, and the precipitate washed and dried over a water-bath), it is the usual and the Pharmacopoeial process for the preparation of "white precipitate," the old "ammonio-chloride" or "amido-chloride of mercury" (so-called because then considered to be a compound of mercury with chlorine and with *amidogen*, NH_2 , or

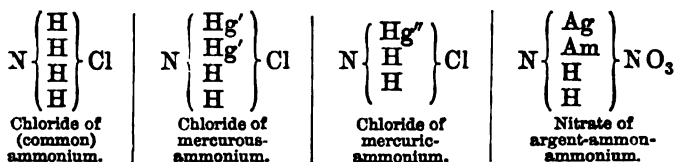
$\text{HgCl}_2 \cdot \text{Hg}_2\text{NH}_3$, chloride and *amide* of mercury), now known as Ammoniated Mercury (*Hydrargyrum Ammoniatum*, B. P.).

Constitution of Ammoniated Mercury.—This precipitate is considered to be the chloride of mercuric-ammonium ($\text{NH}_2\text{Hg}''\text{Cl}$)—that is, chloride of ammonium (NH_4Cl) in which two univalent atoms of hydrogen are displaced by one bivalent atom of mercury.

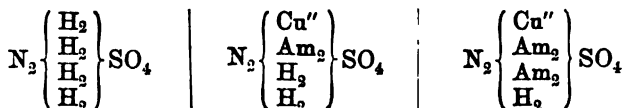


Varieties of Ammoniated Mercury.—If the order of mixing be reversed and ammonia be added to solution of mercuric chloride, a double chloride of mercuric-ammonium and mercury results (NH_2HgCl , HgCl_2): it contains 76.55 per cent. of mercury. Previously to the year 1826, "white precipitate" was officially made by adding a fixed alkali to a solution of equal parts of corrosive sublimate and sal-ammoniac; this gave a double chloride of mercuric-ammonium and ammonium (NH_2HgCl , NH_4Cl), containing 65.57 per cent. of mercury. This compound is now known as "*fusible white precipitate*," because at a temperature somewhat below redness it fuses and then volatilizes. The "white precipitate" which has been official since 1826 contains 79.52 per cent. of mercury. The true compound may be distinguished as "*infusible white precipitate*," from the fact that when heated it volatilizes without fusing. An ointment of this body is official (*Unguentum Hydrargyri Ammoniaci*, B. P.). Prolonged washing with water converts "white precipitate" into a yellowish compound (NH_2HgCl , HgO); hence the official preparation is seldom thoroughly freed from the chloride of ammonium which is formed during its manufacture, and which, if present in larger proportion than seven or eight per cent., gives to it the character of partial or complete fusibility. With iodine, chlorine, or bromine, white precipitate may yield the highly explosive iodide, chloride, or bromide of nitrogen.

Note.—Chloride of mercuric-ammonium is only one member of a large class of compounds derivable from the various salts of ammonium by displacement of atoms of hydrogen in the molecules by atoms of other radicals. The composition of the chloride of mercurous-ammonium (*see* next page) and of ammonio-nitrate of silver is consistent with this view.



The composition of the ammonio-sulphates of copper (pp. 198 and 217) is consistent with the second and third of the following formulæ, the first being that of sulphate of ammonium:—



The iodide of dimercuric-ammonium (NHg''_2I) is formed in testing for ammonia by the "Nessler" reagent (*vide Index*).

Fourth Analytical Reaction.—Pass sulphuretted hydrogen through a mercuric solution; a black precipitate of mercuric sulphide (HgS) falls.

Note.—Sulphuretted hydrogen also precipitates mercurous sulphide (Hg_2S) from mercurous solutions; and in appearance the precipitates are alike; hence this reagent does not distinguish between mercurous and mercuric salts. But in the course of systematic analysis, mercuric salts are thrown down from solution as sulphide after mercurous salts have been otherwise removed. The sulphides are insoluble in sulphhydrate of ammonium.

Note.—An insufficient amount of the gas gives a white or coloured precipitate of oxysulphide. Prolonged contact with sulphuretted hydrogen-water or a sulphhydrate, especially when the mixture is kept warm, converts the black into a red sulphide.

Æthiop's Mineral, the *Hydrargyri Sulphuretum cum Sulphure*, is a mixture of sulphide of mercury and sulphur, obtained on triturating the elements in a mortar till globules are no longer visible. Its name is probably in allusion to its similarity in colour to the skin of the *Æthiop*. It was formerly official.

Vermilion is mercuric sulphide prepared by sublimation.

Tests, continued. (The mercury occurring as mercurous salt.)

Fifth Analytical Reaction.—To a solution of a mercurous salt (the mercurous nitrate obtained in the second synthetical reaction, for example) add hydrochloric acid or any soluble chloride; a white precipitate of calomel (HgCl) occurs.

This reaction was formerly official in the Dublin Pharmacopœia as a process for the preparation of calomel.

Sixth Analytical Reaction.—To solution of a mercurous salt add iodide of potassium; green mercurous iodide (HgI) is precipitated.

Seventh Analytical Reaction.—To a mercurous salt, dissolved or undissolved (e.g. calomel), add ammonia; black salt (e.g. chloride) of mercurous-ammonium ($\text{NH}_2\text{Hg}_2\text{Cl}$) is formed. (See previous page.)

Other Tests for Mercury.

The elimination of mercury in the actual state of metal by the copper test, coupled with the production or non-production of a white precipitate on the addition of hydrochloric acid to the original solution, is usually sufficient evidence of the presence of mercury and its existence as a mercurous or mercuric salt. But other tests may sometimes be applied with advantage. Thus metallic mercury is deposited on placing a drop of the solution on a plate of gold (sovereign or half-sovereign), and touching the drop and the edge of the plate simultaneously with a key; an electric current passes, under these circumstances, from the gold to the key, and thence through the liquid to the gold, decomposing the salt, the mercury of which forms a white metallic spot on the gold, while the other elements go to the iron. This is called *the galvanic test*, and is useful for clinical purposes.—Solution of stannous chloride (SnCl_2)—see Index,—from the readiness with which it forms stannic salts (SnCl_4 , SnO_2 , etc.), gives a white precipitate of mercurous chloride in mercuric solutions, and quickly still further reduces this mercurous chloride (and

other mercury-salts) to a greyish mass of finely divided mercury; this is the old *magpie test*, probably so called from the white and grey appearance of the precipitate. The reaction may even be obtained from such insoluble mercury compounds as "white precipitate."—Confirmatory tests for mercuric and mercurous salts will be found in the action of solution of potash, solution of soda, lime-water, solution of ammonia, and solution of iodide of potassium. (*Vide* pages 230, 232, and 223.)—Normal alkaline carbonates produce yellowish mercurous carbonate and brownish-red mercuric carbonate, both of them unstable.—Alkaline bicarbonates give mercurous carbonate with mercurous solutions, and, with mercuric salts, white (becoming red) mercuric oxysalt.—Yellow chromate of potassium (K_2CrO_4) gives, with mercurous salts, a red precipitate of mercurous chromate (Hg_2CrO_4).—Mercury and all its compounds are volatile, many of them being decomposed at the same time, and yielding globules of condensed metal; the experiment is most conveniently performed in a test-tube.

Antidote.—Albumen gives a white precipitate with solution of mercuric salts; hence the importance of administering white of egg, while waiting for a stomach-pump, in case of poisoning by corrosive sublimate.

QUESTIONS AND EXERCISES.

291. Name the chief ore of mercury, and describe a process for the extraction of the metal.

292. Give the properties of mercury.

293. In what state does mercury exist in "Grey Powder"?

294. What other preparations of metallic mercury itself are employed in medicine?

295. State the relation of the mercurous to the mercuric compounds.

296. Distinguish between an *alloy* and an *amalgam*.

297. State the formulæ of the two Iodides of Mercury.

298. Under what circumstances does mercuric iodide assume two different colours?

299. Illustrate the chemical law of Multiple Proportions as explained by the atomic theory, employing for that purpose the stated composition of the two iodides of mercury.

300. Write down the formulæ of Mercurous and Mercuric Nitrates and Sulphates.

301. How is Mercuric Sulphate prepared?

302. What is the formula of "Turpeth Mineral"?

303. Describe the processes necessary for the conversion of mercury into Calomel and Corrosive Sublimate, using diagrams.

304. Why is black oxide of manganese sometimes mixed with the other ingredients in the preparation of corrosive sublimate?

305. Give the chemical and physical points of difference between calomel and corrosive sublimate.

306. How may a small quantity of calomel in corrosive sublimate be detected?

307. Work out a sum showing how much mercury will be required in the manufacture of one ton of Calomel. *Ans.* 17 cwt. nearly.

308. Mention the official preparations of the chlorides of mercury.

309. Give the formulæ and mode of formation of the Red, Yellow, and Black Oxides of Mercury, employing diagrams.

310. Explain the action of the chief general test for mercury.

311. How are mercurous and mercuric salts analytically distinguished?

312. Give a probable view of the constitution of *Hydrargyrum Ammoniatum*, B. P., and an equation showing how it is made.

313. What is the best temporary antidote in cases of poisoning by mercury?

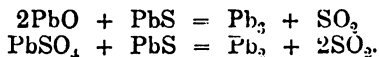
LEAD.

Symbol Pb. Atomic weight 207.

Source.—The ores of lead are numerous; but the one from which the metal is chiefly obtained is the sulphide of lead (PbS), or *galena* (from γαλήνη, *galēnē*, tranquillity, perhaps from its supposed effect in allaying pain).

Preparation.—The ore is first roasted in a current of air; much sulphur is thus burnt off as sulphurous acid gas, while some of the metal is converted into oxide and a portion of the sulphide oxidized to sulphate. Oxidation being stopped when

the mass presents certain appearances, the temperature is raised, and the oxide and sulphate, reacting on undecomposed sulphide, yield the metal and much sulphurous acid gas :—



Uses.—The uses of lead are well known. Alloyed with arsenicum it forms common *shot*, with antimony gives *type-metal*, with tin *solder*, and in smaller quantities enters into the composition of *Britannia metal*, *pewter*, and other alloys. Lead is so slightly attacked by acids that chemical vessels and instruments are often made of it. Even hot hydrochloric acid only slowly converts it into chloride of lead with evolution of hydrogen. Sulphuric acid by aid of air only very slowly attacks it with formation of sulphate of lead and water. Even nitric acid very slowly converts it into nitrate with evolution of nitric oxide and nitrous oxide gases and water.

The salts of lead used in pharmacy and all other preparations of lead are obtained, directly or indirectly, from the metal itself. Heated in a current of air, lead combines with oxygen and forms oxide of lead (PbO), a yellow powder (*massicot*), or, if fused and solidified, a brighter, reddish-yellow heavy mass of bright scales (*Plumbi Oxidum*, B. P.), termed *litharge* (from *λίθος*, *lithos*, a stone, and *ἀργυρος*, *argyros*, silver). It is from this oxide that the chief lead compounds are obtained. Oxide of lead, by further roasting in a current of air, yields *red lead* (or *minium*), Pb_3O_4 , or $\text{PbO}_2 \cdot 2\text{PbO}$. Both oxides are much used by painters, paper-stainers, and glass-manufacturers. *White lead* is a mixture of carbonate (PbCO_3) and hydrate of lead (Pb_2HO) (commonly 2 molecules of the former to 1 of the latter), usually ground up with about 7 per cent of linseed-oil; it is made by exposing lead, cast in spirals or little gratings, to the action of air, acetic fumes, and carbonic acid, the latter generated from decaying vegetable matter, such as spent tan: oxy-acetate of lead slowly but continuously forms, and is as continuously decomposed by the carbonic acid with production of hydrate and carbonate, or *dry white lead*. The grating-like masses, when ground, form the heavy white pulverulent official *Plumbi Carbonas*. The latter is the active constituent of *Unguentum Plumbi Carbonatis*, B. P., the old *Unguentum Cerussæ*.

Lead compounds are poisonous, producing saturnine colic, or even paralysis. These effects are termed *saturnine* from an old name of lead, *Saturn*. The alchemists called lead Saturn, first, because they thought it the oldest of the seven then known metals, and it might therefore be compared to Saturn, who was supposed to be the father of the gods, and, secondly, because its power of dissolving other metals recalled a peculiarity of Saturn, who was said to be in the habit of devouring his own children.

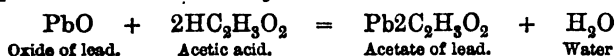
Quantivalence.—The atom of lead is sometimes quadrivalent ($\text{Pb}^{''''}$); but in most of the compounds used in medicine it exerts bivalent activity only ($\text{Pb}^{''}$).

REACTIONS HAVING (a) SYNTHETICAL AND (b) ANALYTICAL INTEREST.

(a) *Synthetical Reactions.*

Acetate of Lead.

First Synthetical Reaction.—Place a few grains of oxide of lead in a test-tube, add about an equal weight of water and two and a half times its weight of acetic acid, and boil; the oxide dissolves (or, rather, disappears—dissolves with simultaneous decomposition) and forms a solution of acetate of lead ($\text{Pb2C}_2\text{H}_3\text{O}_2$). When cold, or on evaporation if much water has been used (the solution being kept faintly acid), crystals of acetate of lead ($\text{Pb2C}_2\text{H}_3\text{O}_2, 3\text{H}_2\text{O}$) are deposited. Larger quantities are obtained by the same method.



This is the official process for *Plumbi Acetas*, B. P. The salt is termed *Sugar of Lead*, from its sweet taste. Besides its direct use in pharmacy, it forms three-fourths of the *Pilula Plumbi cum Opio*, B. P., is the chief constituent of *Unguentum Plumbi Acetatis*, and an ingredient in *Suppositoria Plumbi Composita*.

Subacetate or Oxyacetate of Lead.

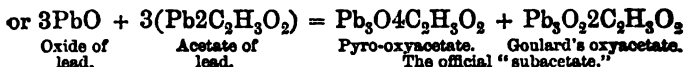
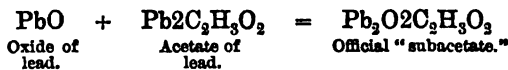
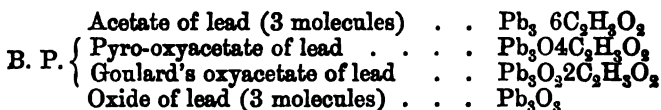
Second Synthetical Reaction.—Boil acetate of lead with about four times its weight of water and rather more than

two-thirds its weight of oxide of lead; the resulting filtered liquor is solution of oxyacetate of lead, *Liquor Plumbi Subacetatis*, B. P.

The official *Liquor* is made by boiling 5 ounces of acetate and $3\frac{1}{2}$ of oxide in one pint of distilled water for half an hour (constantly stirring), filtering, and making up for any loss during evaporation by diluting the filtrate to 1 pint.

A similar solution was used by M. Goulard, who called it *Extractum Saturni*, and drew attention to it in 1770. It is now frequently termed *Goulard's Extract*. A more dilute solution, 1 of *Liquor* and 1 of spirit in 80 of distilled water, is also official in the British Pharmacopœia, under the name of *Liquor Plumbi Subacetatis Dilutus*. The latter is commonly known as *Goulard Water*. The stronger solution is the chief ingredient in *Unguentum Plumbi Subacetatis Compositum*, B. P., a slight modification of the old *Goulard's Cerate*.

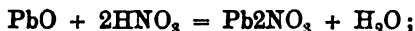
Oxyacetates of lead.—The official subacetate of lead is not a definite chemical salt. It is probably a mixture of two subacetates of lead, which are well-known crystalline compounds, and which the author is disposed to regard as having a constitution similar to that he has already indicated for some other salts (*see* Iron and Antimony, also Bismuth). Exposed to air it absorbs carbonic acid gas, and hydrato-carbonate of lead is deposited.



Nitrate of Lead. Red Lead. Peroxide of Lead.

Third Synthetical Reaction.—Digest a few grains of red lead in nitric acid and water; nitrate of lead (Pb2NO_3) is formed, and remains in solution, while a puce-coloured peroxide of lead (PbO_2) is precipitated.

Nitrate of lead (*Plumbi Nitras*, B. P.) is more directly made by dissolving litharge (PbO) in nitric acid,

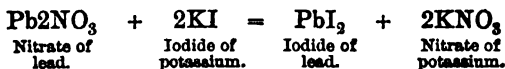


but the former reaction serves to bring before the reader two other oxides of lead, namely red lead (Pb_3O_4) and peroxide of lead (PbO_2). In the latter oxide the quadrivalent character of lead is obvious. Nitrate of lead is used officially in preparing iodide of lead; for this purpose the above mixture is filtered, the precipitate of peroxide of lead purified from adhering nitrate by passing hot water through the filter, the filtrate and washings evaporated to dryness to remove excess of nitric acid, the residual nitrate of lead redissolved by ebullition with a small quantity of hot water, and the solution set aside to crystallize, or a portion at once used for the following experiment. Nitrate of lead forms white crystals derived from octahedra.

Peroxide of lead dissolved in strong hydrochloric acid apparently yields an unstable perchloride (PbCl_4).

Iodide of Lead.

Fourth Synthetical Reaction.—To a neutral solution of nitrate of lead add solution of iodide of potassium; a precipitate of iodide of lead (PbI_2) falls (*Plumbi Iodidum*, B. P.). Equal weights of the salts may be used in making large quantities.

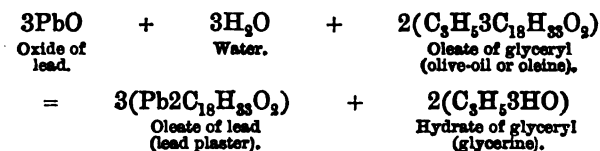


Iodide of lead is the chief ingredient in *Emplastrum Plumbi Iodidi*, B. P.; and *Unguentum Plumbi Iodidi*, B. P. The iodide of lead used in making the plaster is altered in the process, either iodide of sodium being formed (from the hard soap present), or possibly a compound of the iodide and oleate of lead, corresponding with the almost colourless hydrato-iodides of lead.

Crystals of Iodide of Lead.—Heat the iodide of lead with the supernatant liquid, and, if necessary, filter; the salt is dissolved, and again separates in golden crystalline scales as the solution cools.

Oleate of Lead. (Lead Plaster.)

Fifth Synthetical Reaction.—Boil together in a small dish a few grains of very finely powdered oxide of lead, rather more than twice its weight of olive-oil, and ten or twenty times as much water, well stirring the mixture, and from time to time replacing water that has evaporated; the product is a white mass of oleate of lead ($\text{Pb}_2\text{C}_{18}\text{H}_{33}\text{O}_2$) (*Emplastrum Plumbi*, B.P.), glycerine remaining in solution in the water. Larger quantities are prepared in the same manner.



The action between the oxide of lead and olive-oil is slow, requiring several hours for its completion. Lead plaster is a constituent of eleven of the fourteen plasters (*Emplastra*) mentioned in the British Pharmacopœia.

The glycerine may be obtained by treating the aqueous product of the above reaction with sulphuretted hydrogen to remove a trace of lead, then digesting with animal charcoal, filtering, and evaporating. But on the large scale glycerine is produced as a by-product in the manufacture of candles; for its elements are found in nearly all vegetable and animal fats. (*Vide Index*.) If in making lead plaster the mixture be evaporated to dryness, some of the glycerine will escape with the steam and some remain with the plaster.

Modes of formation of *Chloride, Sulphide, Chromate, Sulphate, Hydrate*, and other salts of lead are incidentally described in the following analytical paragraphs.

(b) Reactions having Analytical Interest (Tests).

First Analytical Reaction.—To a solution of lead salt (acetate, for example) add hydrochloric acid; a white precipitate of chloride of lead (PbCl_2) is obtained. Boil the precipitate with much water; it dissolves, but, on the solution cooling, is redeposited in small acicular crystals. Filter the cold solution, and pass sulphuretted hydrogen through it; a black

precipitate (sulphide of lead, PbS) shows that the chloride of lead is soluble to a slight extent in cold water.

Note.—A white precipitate on the addition of hydrochloric acid, soluble in hot water, and blackened by sulphuretted hydrogen, sufficiently distinguishes lead salts from those of other metals; but the non-production of such a precipitate does not prove the absence of a small quantity of lead, chloride of lead being slightly soluble in cold water. Hydrochloric acid will be found to be a useful but not a delicate test for lead.

Second Analytical Reaction.—Through a dilute solution of a lead salt, acidulated with hydrochloric acid, pass sulphuretted hydrogen; a black precipitate of sulphide of lead (PbS) occurs.

Lead in Water.—The foregoing is a very delicate test. Should a trace of lead be present in water used for drinking purposes, sulphuretted hydrogen will detect it. On passing the gas through a pint of such (acidulated) water, a brownish colour is produced. If the tint is scarcely perceptible, set the liquid aside for a day; the gas will become decomposed and a thin layer of sulphur be found at the bottom of the vessel, white if no lead be present, but more or less brown if it contain sulphide of lead.

Third Analytical Reaction.—To solution of a lead salt add sulphhydrate of ammonium; a black precipitate of sulphide of lead falls, insoluble in excess.

Fourth Analytical Reaction.—To solution of a lead salt add solution of chromate of potassium (K_2CrO_4); a yellow precipitate of chromate of lead ($PbCrO_4$) is formed, insoluble in weak acids.

Chromes.—This reaction has technical as well as analytical interest. The precipitate is the common pigment termed *chrome yellow*, or *lemon chrome*. Boiled with lime and water, a portion of the chromic radical is removed as a soluble chromate of calcium, and an oxychromate of lead, of a bright red or orange colour (*orange chrome*), is produced.

Fifth Analytical Reaction.—To solution of a lead salt add

dilute sulphuric acid, or solution of a sulphate; a white precipitate of sulphate of lead (PbSO_4) falls.

Sulphate of lead is slightly soluble in strong acids, and in solutions of alkaline salts; it is insoluble in acetic acid. It is readily dissolved and, indeed, decomposed by solution of acetate of ammonium, the liquid yielding the ordinary reactions with soluble chromates and iodides.

In dilute solutions the above sulphuric reaction does not take place immediately; the precipitate, however, falls after a time; its appearance may be hastened by evaporating the mixture nearly to dryness and then rediluting.

The white precipitate always noticed in the vessels in which diluted sulphuric acid is kept is sulphate of lead, derived from the leaden chambers in which the acid is made: solubility in strong acid and insolubility in weak explains its appearance.

Antidotes.—From the insolubility of sulphate of lead in water, the best *antidote* in a case of poisoning by the acetate or other soluble salt of lead is a soluble sulphate, such as Epsom salt, sulphate of sodium, or alum, vomiting being also induced or the stomach-pump applied as quickly as possible.

Other tests for lead will be found in the reaction with *iodide of potassium* (*vide* p. 241); with *alkaline carbonates*, a white precipitate ($2\text{PbCO}_3 + \text{Pb}_2\text{HO}$) insoluble in excess; with *alkalies*, a white precipitate (Pb_2HO) more or less soluble in excess; with *alkaline phosphates, arseniates, ferrocyanides and cyanides*, precipitates mostly insoluble, but of no special analytical interest. Insoluble salts of lead are decomposed by solutions of potash (KHO) or soda (NaHO).

The metal is precipitated in a beautifully crystalline state by metallic zinc and some other metals; the *lead tree* is thus formed.—The *blowpipe-flame* decomposes solid lead compounds placed in a small cavity in a piece of charcoal, a soft malleable bead of metal being produced, and a yellowish ring of oxide deposited on the charcoal.

QUESTIONS AND EXERCISES.

314. Write down equations descriptive of the smelting of galena.

315. Mention some of the alloys of lead.
316. How is litharge produced?
317. Give the formulæ of white lead and red lead.
318. Describe the manufacture of white lead.
319. What is the quantivalence of lead?
320. Draw a diagram expressive of the formation of ordinary Acetate of Lead.
321. Describe the preparation and composition of *Liquor Plumbi Subacetatis*, B. P.
322. What is the action of nitric acid on red lead, litharge, and metallic lead?
323. How is the official Iodide of Lead prepared?
324. Describe the reaction between oxide of lead, water, and olive-oil, at the temperature of boiling water, and give chemical formulæ explanatory of the constitution of the products.
325. Mention the chief tests for lead.
326. How would you search for lead in potable water?
327. What is the composition of chrome yellow?
328. State a method whereby lead, barium, and silver may be separated from each other.
329. Name the best antidote in cases of poisoning by the soluble salts of lead.

SILVER.

Symbol Ag. Atomic weight 108.

Source.—This element occurs in nature in the free state and as ore, the common variety of the latter being sulphide of silver (Ag_2S) in combination with much sulphide of lead, forming *argentiferous galena*.

Preparation.—The lead from such galena (p. 237) is melted and slowly cooled; crystals of lead separate and are raked out from the still fluid mass, and thus an alloy very rich in silver is finally obtained; this is roasted in a current of air, whereby the lead is oxidized and removed as litharge, pure silver remaining. Other ores undergo various preparatory treatments according to their nature, and are then shaken with mercury, which amalgamates with and dissolves the particles of silver, the mercury being subsequently removed from the amalgam by distillation. Soils and minerals containing metallic silver are also treated in this way. An important improvement in the amalgamation process, by which

the mercury more readily unites with the silver, consists in the addition of a small proportion of sodium to the mercury—a discovery simultaneously made in England by Crookes and in New York by Wurtz.

Silver is not readily affected by the weak acids or other fluids of food, though it is rapidly tarnished by sulphur or sulphur compounds. It does not perceptibly attack hydrochloric acid; reduces strong nitric acid to nitrous anhydride (N_2O_3), and a weaker acid to nitric oxide (NO); it reduces hot sulphuric acid to sulphurous anhydride (SO_2), sulphate of silver (Ag_2SO_4) being formed. The latter salt is crystalline, and slightly soluble in water.

REACTIONS HAVING (a) SYNTHETICAL AND (b) ANALYTICAL INTEREST.

(a) *Synthetical Reactions.*

Impure Nitrate of Silver.

First Synthetical Reaction.—Dissolve a silver coin in nitric acid; nitric oxide gas (NO) and nitrous anhydride (N_2O_3) are evolved, and a solution of nitrates of silver and copper is obtained.

Silver Coinage.—Pure silver is too soft for use as coin; it is therefore hardened by alloying with copper. The silver money of England contains 7·5, of Prussia 25, and of France 10 and 16·5 per cent. of copper; for the fineness of the French standard silver is 0·900 in the five-franc piece, while an inferior alloy of 0·835 is used for the lower denominations. The single-franc piece, composed of the latter alloy, is still made to weigh five grammes, the weight originally chosen for the franc as the unit of the monetary scale when the fineness of the coin was 0·900. It has now become a token, like the British shilling, of which the nominal value exceeds the metallic value. One pound troy of British standard silver is coined into 66 shillings, of which the metal is worth from 60s. to 62s. according to the market price of silver. The standard fineness of this silver is 0·925, three alloy in 40. British silver coins are a legal tender in payments to the amount of 40s. only

Chloride of Silver.

Second Synthetical Reaction.—To the product of the foregoing reaction add water and hydrochloric acid or a soluble chloride; white chloride of silver (AgCl) is precipitated, copper still remaining in solution. Collect the precipitate on a filter and wash with water; it is pure chloride of silver.

Note.—The nitrates of silver and copper may also be separated by evaporating the solution of the metals in nitric acid to dryness and gently heating the residue, when the nitrate of copper is decomposed but the nitrate of silver is unaffected. The latter may be dissolved from the residual oxide of copper by water.

Chloride of silver may be obtained in crystals by evaporation of its solution in ammonia.

Pure Silver.

Third Synthetical Reaction.—Place the chloride of silver of the previous reaction in a dish, wet it with dilute sulphuric acid, and float a piece of sheet zinc on the mixture; metallic silver is precipitated, and after about one day wholly removed from solution. Collect the precipitate on a filter and wash with water; it is pure metallic silver, and is readily fusible into a single button.

Note.—Any considerable quantity of chloride of silver may also be reduced to a lump of the metal by fusion, in a crucible, with about half its weight of carbonate of sodium.

Pure Nitrate of Silver.

Fourth Synthetical Reaction.—Dissolve the pure silver of the previous reaction in nitric acid (3 of silver require about 2 or $2\frac{1}{2}$ of strong acid diluted with 5 of water), and remove excess of acid by evaporating the solution to dryness, slightly heating the residue; the product is pure nitrate of silver. Dissolve by heating with a small quantity of water; on the solution cooling, or on evaporation, colourless tabular crystals of nitrate of silver are obtained.



Notes.—The solution of pure or refined silver (*Argentum Purificatum*, B. P.) in nitric acid, evaporation, and crystallization constitute the official process for the preparation of the nitrate (*Argenti Nitras*, B. P.). The salt fused, and poured into proper moulds, yields the white cylindrical sticks or rods commonly termed *caustic* (from *kaiw*, *kaiō*, I burn), or *lunar caustic*. (The alchemists called silver *Diana* or *Luna*, from its supposed mysterious connection with the moon.) The specimen of nitrate of silver obtained in the foregoing reaction, dissolved in water, will be found useful as an analytical reagent. Nitrate of silver is soluble in rectified spirit; but after a time reaction and decomposition occur.

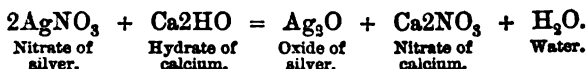
Silver salts are decomposed when in contact with organic matter, especially in the presence of light or heat, the metal itself being liberated, or a black insoluble compound formed. Hence the value of the nitrate in the manufacture of indelible ink for marking linen; hence, too, the reason of the practice of rendering silver solutions clear by subsidence and decantation, rather than by filtration through paper; and hence the cause of those cases of actual combustion which have been known to occur in preparing pills containing oxide of silver and essential oil or other organic matter. Linen marked with such ink should not be cleansed by aid of bleaching liquor, as the marked parts are then apt rapidly to be oxidized into perfectly rotten matter, holes resulting. Paul says the reaction is as follows:— $\text{Ag}_2\text{O} + \text{CaCl}_2\text{O}_2 = 2\text{AgCl} + \text{CaO} + \text{O}_2$.

Oxide of Silver.

Fifth Synthetical Reaction.—To a few drops of solution of nitrate of silver add solution of potash or soda or lime-water; an olive-brown precipitate of oxide of silver (Ag_2O) occurs. The washed and dry oxide, like most silver compounds, is decomposed by heat with production of metal. It is also reduced by contact with organic matter. (*See the previous paragraph.*)

The *Argenti Oxidum*, B. P., is thus made, lime-water being the precipitant employed, soda or potash not being so readily

removed by washing. Three and a half pints of good lime-water will decompose half an ounce of nitrate of silver.



Methods of forming several other salts of silver are incidentally mentioned in the following analytical paragraphs.

(b) *Reactions having Analytical Interest (Tests).*

First Analytical Reaction.—To a solution of a silver salt add hydrochloric acid or other soluble chloride; a white curdy precipitate of chloride of silver falls. Add nitric acid, and boil; the precipitate does not dissolve. Pour off the acid and add solution of ammonia; the precipitate dissolves. Neutralize the ammoniacal solution by an acid; the chloride of silver is reprecipitated.

This is the most characteristic test for silver. The precipitated chloride is also soluble in solutions of hyposulphite of sodium or cyanide of potassium—facts of considerable importance in photographic operations.

Other analytical reagents than the above are occasionally useful.—Sulphuretted hydrogen, or sulphhydrate of ammonium, gives a black precipitate, sulphide of silver (Ag_2S), insoluble in alkalies.—Solutions of potash or soda give a brown precipitate, oxide of silver (Ag_2O), converted into a fulminating compound by prolonged contact with ammonia.—Phosphate of sodium gives a pale yellow precipitate, phosphate of silver (Ag_3PO_4), soluble in nitric acid and in ammonia.—Arsenate of ammonium gives a chocolate-coloured precipitate, arseniate of silver (Ag_3AsO_4), already noticed in connection with arsenic acid.—Iodide or bromide of potassium gives a yellowish white precipitate, iodide or bromide of silver (AgI or AgBr), insoluble in acids and only slightly soluble in ammonia.—Cyanide of potassium gives a white precipitate, cyanide of silver (AgCy), soluble in excess,

sparingly soluble in ammonia, insoluble in dilute nitric acid, soluble in boiling concentrated nitric acid.——Yellow chromate of potassium (K_2CrO_4) gives a red precipitate, chromate of silver (Ag_2CrO_4).——Red chromate of potassium also gives a red precipitate, acid chromate of silver (Ag_2CrO_4 , CrO_3).——Many organic acids afford insoluble salts of silver.——Several metals displace silver from solution, mercury forming in this way a crystalline compound known as the silver tree, or *Arbor Dianæ*.——In the blowpipe-flame, silver salts, placed on charcoal with a little carbonate of sodium, yield bright globules of metal, accompanied by no incrustation as in the corresponding reaction with lead salts; the experiment may be performed with the nitrate, which first melts and then, like all nitrates, deflagrates, yielding a white metallic coating of silver which slowly aggregates to a button.

Antidotes.—Solution of common salt, sal-ammoniac, or any other inert chloride should obviously be administered where large doses of nitrate of silver have been swallowed. A quantity of sea-water or brine would convert the silver into insoluble chloride, and at the same time produce vomiting.

DIRECTIONS FOR APPLYING SOME OF THE FOREGOING REACTIONS TO THE ANALYSIS OF AN AQUEOUS SOLUTION OF A SALT OF ONE OF THE METALS COPPER, MERCURY (EITHER AS MERCUROUS OR MERCURIC SALT), LEAD, SILVER.

Add hydrochloric acid :—

Silver is indicated by a white curdy precipitate, soluble in ammonia.

Mercurous salts also by a white precipitate, turned black by ammonia.

Lead by a white precipitate, insoluble in ammonia. Confirm by boiling another portion of the hydrochloric precipitate in water; it dissolves.

If hydrochloric acid gives no precipitate, silver and mercurous salts are absent. Lead can only be present in very small

quantity. Mercuric salts may be present. Copper may be present. Divide the liquid into three portions, and apply a direct test for each metal, as follows.

Lead is best detected by the sulphuric test, the tube being set aside for a time if the precipitate does not appear at once.

Mercury is best detected by the copper test. If present here, it occurs as mercuric salt.

Copper betrays itself by the blue colour of the liquid under examination. Confirm by the ammonia test.

If the above reactions are not thoroughly conclusive, confirmatory evidence should be obtained by the application of some of the other reagents for copper, mercury, lead, or silver.

TABLE OF SHORT DIRECTIONS FOR APPLYING SOME OF THE FOREGOING REACTIONS TO THE ANALYSIS OF AN AQUEOUS SOLUTION OF SALTS OF ANY OR ALL OF THE METALS COPPER, MERCURY (EITHER MERCUROUS OR MERCURIC SALT, OR BOTH), LEAD, SILVER.

Add hydrochloric acid, filter, and wash the precipitate with a small quantity of cold water.

Ppt. Pb Hg (ous) Ag. Wash with boiling water.		Filtrate Cu Hg (ic) Pb. Divide into three portions. Test for		
Ppt. Hg (ous) Ag. Add AmHO.	Filtrate Pb. Add H ₂ SO ₄ , white ppt.*	Cu by AmHO; blue sol. Hg (mercuric) by Cu; globules. Pb by H ₂ SO ₄ ; white ppt.*		
Precipitate Hg (mercurous), black.	Filtrate Ag. Add HNO ₃ , white ppt.			

* Liquids containing only a small quantity of lead do not readily yield sulphate of lead on the addition of sulphuric acid. Before lead can be said to be absent, therefore, the liquid should be evaporated to dryness with one drop of sulphuric acid, and the residue digested in water; any sulphate of lead then remains as a heavy white insoluble powder.

SHORT DIRECTIONS FOR THE ANALYSIS OF AN AQUEOUS SOLUTION OF ORDINARY SALTS OF ONE
OF THE ELEMENTS HITHERTO CONSIDERED.

Add hydrochloric acid.

Ppt. Hg(ous) Pb Ag. Add AmHO. Hg(ous), black ppt. Pb, ppt. still white. Ag, ppt. dissolved. <i>Note I.</i>—If HCl gave no precipitate, neither Hg(ous) nor Ag is present; and Pb only in minute amount, if at all. <i>Note II.</i>—Hg obtained here must have existed in the solution as a mercurous salt. <i>Note III.</i>—Sb is also precipitated by HCl, but is dissolved on adding more HCl; the Hg, Pb, and Ag precipitates are not soluble in excess of HCl.	Ppt. Cu Hg(ic) Pb As Sb. As, yellow ppt. Sb, orange ppt. Cu Hg(ic) } black ppt. Pb Test original solution for Cu by AmHO; blue sol. Hg by Cu; globules. Pb by H₂SO₄; white ppt. <i>Note I.</i>—If H₂S gave no precipitate, neither Cu, Hg, Pb, As, nor Sb is present. <i>Note II.</i>—Hg and Pb may give coloured precipitates (oxysulphides) with H₂S if too little of the latter has been passed through the solution.	If HCl gave no precipitate the metal is still in the liquid; pass H₂S through it. Ppt. Fe Al Zn. Fe, black ppt. Test original solution for ferric salt by K₄Fcy (dark blue ppt.); and for ferrous salt by K₆Fdcy (dark blue ppt.). Al } white ppt. Zn } Test original solution by AmHO. Al, white ppt. insoluble in excess. Zn, white ppt. soluble in excess.	If H₂S gave no precipitate the metal is still in the liquid; add AmCl, AmHO, and AmHS. If AmHS, etc. gave no precipitate, the liquid may still contain either Ba, Ca, Mg, K, Na, or Am; add successively K₂CrO₄ for Ba, Am₂C₂O₄ for Ca, Na₂HPO₄ for Mg. If neither Ba, Ca, nor Mg is found, examine the original solution for Am by KHO, Na by the flame test, and K by PtCl₄.
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TABLE OF SHORT DIRECTIONS FOR APPLYING SOME OF THE FOREGOING ANALYTICAL REACTIONS TO THE ANALYSIS OF AN AQUEOUS SOLUTION OF ORDINARY SALTS OF ANY OR ALL OF THE ELEMENTS HITHERTO CONSIDERED.

(READ THE "MEMORANDA" ON THE FOLLOWING PAGE.)

Add hydrochloric acid, and filter.

Precipitate Hg(ous) Pb Ag. Wash, boil with water, filter.	Precipitate Hg(ous) Ag. Add AmHO to the well washed precipitate on the filter.	Filtrate Pb. Add H ₂ SO ₄ ; white precip- itate.	Precipitate Cu Hg(ce) Pb As Sb. Wash, digest in AmHS, filter.	Filtrate As Sb. Add HCl, H ₂ O, and boil; digest the precipitate in strong HCl; boil, dilute; filter.	Precipitate Fe Al Zn. Wash, dissolve in HCl, boil (with a few drops of HNO ₃ , if necessary; p. 133); add KHO, stir, filter.	Filtrate Ba Ca Mg K Na Am. Add Am ₂ CO ₃ ; boil, filter.	Precipitate Ba Ca. Wash, dissolve in HCl, H ₂ O, boil; add K ₂ CrO ₄ , filter.	Filtrate Ca. Add Am ₂ CO ₃ ; white pre- cipitate.	Ppt. Mg; K Na Am. Evap., ignite, dissol. in H ₂ O; test for K by PtCl ₄ ; yel. ppt. Na by flame; yellow, Am. in orig. sol. by NaHO.
Precip. Hg(ous). Black pre- cipitate. white precip- itate.	Filtrate Ag. Add HNO ₃ ; p. 231.)	Precipitate Cu Hg(ce) Pb As Sb. Wash, dissolve in a few drops of HNO ₃ , and HCl; evap. nearly to dry- ness; re-dis- solve in H ₂ O, divide into three, and test for Cu by Am HO; blue solution. Hg by Cu; globules. Pb by H ₂ SO ₄ ; white precip.	Precip. As. Yellow. Add H ₂ S. Orange precip- itate. Confirm by testing original solution by the H tests.	Precipitate Fe. ? Ferric or ferrous. Test origi- nal solution by yellow and red prussiate.	Filtrate. Al Zn. Neutralize by HCl; and AmHO, stir, filter.	Precip. Ba. Yellow. Add Am ₂ CO ₃ ; white pre- cipitate.	Filtrate Ca. Add Am ₂ CO ₃ ; white pre- cipitate.	Ppt. Mg; K Na Am. Evap., ignite, dissol. in H ₂ O; test for K by PtCl ₄ ; yel. ppt. Na by flame; yellow, Am. in orig. sol. by NaHO.	

OUTLINE OF THE PRECEDING TABLES.

H Cl	H ₂ S	AmHS	Am ₂ CO ₃	Am ₂ HA ₂ O ₄	
Hg (as mercurous salt)	Cu	Zn	Ba	Mg	K
Pb (partially)	Hg (as mercuric salt)	Al	Ca		Na
Ag	Pb (entirely)	Fe			Am
	As				
	Sb				

The practical student should examine solutions containing these common metals until he is able to analyse with facility and accuracy. In this way he will best perceive the peculiarities of each element and their general relations to each other. As the rarer metals are not included here, the tables are not complete analytical schemes; only general memoranda respecting them will, therefore, now be given. For *special* memoranda, see p. 298.

MEMORANDA RELATING TO THE GENERAL ANALYTICAL TABLE
(PAGE 253).

The group-tests adopted in the Table are, obviously, *hydrochloric acid, sulphuretted hydrogen, sulphhydrate of ammonium, carbonate of ammonium, and arseniate of ammonium*. If a group-test produces no precipitate, it is self-evident that there can be no member of the group present. At first, therefore, add only a small quantity of a group-test, and if it produces no effect add no more; for it is not advisable to overload a solution with useless reagents; substances expected to come down as precipitates are not unfrequently held in the liquid by excess of acid, alkali, or strong aqueous solution of some group-reagent, thoughtlessly added. Indeed experienced manipulators not unfrequently make preliminary trials with group-reagents on a few drops only of the liquid under examination; if a precipitate is produced, it is added to the

bulk of the original liquid and the addition of the group-reagent continued; if a precipitate is not produced, the few drops are thrown away, and the unnecessary addition of a group-reagent thus avoided altogether, an advantage fully making up for the extra trouble of making a preliminary trial.—While shunning excess, however, care must be taken to avoid deficiency; a substance only partially removed from solution through the addition of an insufficient amount of a reagent will appear where not expected, be constantly mistaken for something else, and cause much trouble; this will not occur if the appearance, odour, or reaction of the liquid on test-paper be duly observed. It is also a good plan, when a group-reagent has produced a precipitate and the latter has been filtered out, to add a little more of the reagent to the clear filtrate; if more precipitate is produced, an insufficient amount of the group-test was introduced in the first instance; but the error is corrected by simply refiltering; if no precipitate occurs, the mind is satisfied and the way cleared for further operations.

Group-precipitates, or any precipitates still requiring examination, should, as a rule, be well washed before further testing; this is to remove the aqueous solution of other substances adhering to the precipitate (the mother-liquor as it is termed), so that subsequent reactions may take place fairly between the reagent used and the precipitate only.—A precipitate is sometimes in so fine a state of division as to retard filtration by clogging the pores of the paper, or even to pass through the filter altogether; in these cases the mixture may be warmed or boiled (or a fresh quantity of the original solution may be warmed *before* the group-test is added), which usually causes aggregation of the particles of a precipitate, and hence facilitates the passage of liquids.

Division of work.—It is immaterial whether a solution be first divided into group-precipitates or each precipitate be examined as soon as produced; if the former method be adopted, confusion will be avoided by labelling or marking the funnels or papers holding the precipitate "the HCl ppt.," "the H_2S ppt.," and so on.

The colours and general appearance of the various sulphides and hydrates precipitated should be borne in mind, as the absence of other bodies, as well as the presence of those thrown down, is often at once thus indicated.

Application of confirmatory tests must be frequent.

Results of analyses should be recorded neatly in a memorandum book.

The various reactions which occur in an analysis have already come before the reader in going through the tests for the individual metals or in other analytical operations; it is unnecessary, therefore, again to draw out equations or diagrams. But the reactions should be thought over, and, if not perfectly clear to the mind, be written out again and again till thoroughly understood.

QUESTIONS AND EXERCISES.

330. By what process is silver obtained from argentiferous galena?

331. What weight of English silver coin will yield one pound of pure nitrate of silver?

332. How may the metal be recovered from an impure mixture of silver salts?

333. Give a diagram showing the formation of nitrate of silver from the metal.

334. Describe the reaction of lime-water and nitrate of silver.

335. Mention the chief test for silver, and the precautions to be observed in order that silver salts may be distinguished from those of lead and mercury.

336. Name the antidote for silver.

337. Give processes for the qualitative analysis of liquids containing the following substances:—

a. Antimony and Mercurous salt.

b. Lead and Calcium.

c. Silver and Mercurous salt.

d. Lead and Mercuric salt.

e. Copper and Arsenicum.

f. Arsenicum and Antimony.

g. Aluminium and Zinc.

h. Iron and Copper.

i. Magnesium, Calcium, and Potassium.

j. Silver, Antimony, Zinc, Barium, and Ammonium.

338. Enumerate the so-called group-tests.
 339. Give a general sketch of the method of analysing a solution suspected to contain two or more salts of common metals.
 340. Classify the common metals according to their analytical relations.

METALS OF MINOR PHARMACEUTICAL IMPORTANCE.

Thus far has been considered, somewhat in detail, the chemistry of the common metals, salts of which are frequently used in medicine or in testing medicinal substances. These are:—

Potassium,	Barium,	Zinc,	Arsenicum,	Mercury,
Sodium,	Calcium,	Aluminium,	Antimony,	Lead,
Ammonium (?),	Magnesium,	Iron,	Copper,	Silver.

Of the remaining metals, nine are mentioned in the British Pharmacopœia, namely—

Lithium,	Chromium,	Cadmium,
Cerium,	Tin,	Platinum,
Manganese.	Gold,	Bismuth.

Compounds of three more are sufficiently common to occasionally come under notice:—

Strontium,	Cobalt,	Nickel
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These twelve metals of minor pharmaceutical interest may be shortly studied, a few only of the reactions of each (just those mentioned in the following pages) being performed. When all have been thus treated, their respective positions in the analytical groups will be indicated and a tabular scheme be given by which an analysis of a solution containing any metal may be effected. Thus, step by step; we may learn how to analyse almost any substance that may occur, and know to what extent the presence of a rarer will interfere with the ordinary tests for a common element; additional illustrations of the working of chemical laws will be acquired, and the store of chemical and pharmaceutical facts be increased. The opportunity thus afforded for improvement is

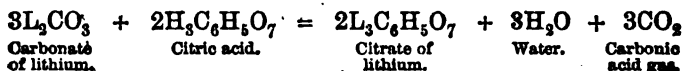
habits of neatness of manipulation, in precision, and in power of classification furnishes another and no mean reason why such experiments should be prosecuted, the direct value of which may not be considerable to medical and pharmaceutical learners.

LITHIUM.

Symbol L. Atomic weight 7.

Lithium is widely distributed in nature, but usually in minute proportions compared with other elements. A trace of it may be found in most soils and waters, a Cornish spring containing even considerable quantities as chloride.

The salt chiefly used in medicine is the *Citrate* ($\text{L}_3\text{C}_6\text{H}_5\text{O}_7$) (*Lithiæ Citras*, B. P.), occurring in white deliquescent crystals or powder, prepared by dissolving 50 grains of the *Carbonate* (L_2CO_3) and 90 of citric acid (50 to 95 if both are quite pure) in 1 ounce of water, evaporating to a low bulk and setting aside in a dry place to crystallize, or at once evaporating to dryness and powdering the residue. The crystals have the formula $\text{L}_3\text{C}_6\text{H}_5\text{O}_7, 4\text{H}_2\text{O}$; dried at 212°F ., $\text{L}_3\text{C}_6\text{H}_5\text{O}_7, \text{H}_2\text{O}$ (Umney).



The carbonate (*Lithiæ Carbonas*, B. P.) is a white granular powder obtained from the minerals which contain lithium—namely, lepidolite (from *λεπίς*, *lepis*, a scale, and *λίθος*, *lithos*, a stone; it has a scaly appearance), triphane (from *τρεῖς*, *treis*, three, and *φαίνω*, *phainō*, I shine) or spodumene (from *σποδῶω*, *spoddōō*, I reduce to ashes, in allusion to its exfoliation in the blowpipe-flame), and petalite (from *πέταλον*, *petalon*, a leaf; its character is leafy and laminated). Each contains silicate of aluminium, with fluoride of potassium and lithium in the case of lepidolite, and silicate of sodium and lithium in the others. *Liquor Lithiæ Effervescens*, B. P., is a solution of 10 grains of carbonate of lithium in 1 pint of water charged with seven times its volume of carbonic acid gas and kept in ordinary aerated water-bottles. "Half a pint, evaporated to dryness, yields 5 grains of a white solid residue, answering to the tests for carbonate of lithium. . . . Ten grains of the latter salt neutralized with sulphuric acid, and afterwards

heated to redness, leaves 14.86 grains of dry sulphate of lithium, which, when redissolved in distilled water, yields no precipitate with oxalate of ammonium or solution of lime," indicating absence of salts of calcium and aluminium. Citrate of lithium should yield by incineration 52.8 per cent. of white carbonate of lithium.

*Urate of lithium** is more soluble than urate of sodium; hence lithium preparations are administered to gouty patients in the hope that urate of sodium, with which such systems are loaded, may be converted into urate of lithium and removed.

In chemical position lithium stands between the alkaline and the alkaline-earth metals, its hydrate, carbonate, and phosphate being slightly soluble in water. The double chloride of platinum and lithium also is soluble in water. Its atom is univalent, L'.

Analytical Reaction.—Moisten the end of a platinum wire with solution of a minute particle of solid lithium salt, and introduce it into the flame of a Bunsen burner or other almost colourless flame (spirit lamp or blowpipe-flame); a magnificent crimson tinge is imparted.

The light thus emitted by ignited lithium vapour is of a purer scarlet than that given by strontium, the next element. When the flames are examined by spectral analysis (physically analysed by a prism), the red rays are, in the case of strontium, found to be associated with blue and yellow, neither of which is present in the lithium light, blue lithium rays only appearing at temperatures much higher than those of the air-gas flame, or, indeed, any other ordinary flame.

STRONTIUM.

Symbol Sr. Atomic weight 87.5.

Source.—Strontium is not widely distributed in nature; but the carbonate (SrCO_3), known as *strontianite*, and the sulphate (SrSO_4), known as *celestine* (from *caelum*, the sky, in allusion to its occasional bluish colour), are by no means rare minerals.

* Urates will be considered subsequently in connection with uric acid.

Salts of strontium are not employed in medicine. They are chiefly used by firework-manufacturers in preparing red fire. The colour they impart to flame is a beautiful crimson—ignited strontium vapour emitting red rays, as already explained. Nitrate of strontium (Sr_2NO_3) is best for pyrotechnic compositions, its oxygen enabling it to burn freely when mixed with charcoal, sulphur, etc. It, or any salts, may be obtained by dissolving the carbonate in the appropriate acid, or by igniting the cheaper sulphate with coal, whereby sulphide (SrS) is produced, and dissolving this in acid.

The position of strontium among the chemical elements is between barium and calcium; its sulphate is extremely sparingly soluble in water. Its atom, like those of barium and calcium, is bivalent (Sr'').

Analytical Reactions (Tests).

First Analytical Reaction.—To solution of a strontium salt (Sr_2NO_3 or SrCl_2) add carbonate of ammonium; a white precipitate of carbonate of strontium (SrCO_3) falls.

Second Analytical Reaction.—To a solution of a strontium salt add sulphuric acid previously so diluted that it will not precipitate calcium salts, or an equally dilute solution of any sulphate, *e.g.* that of calcium itself; a white precipitate of sulphate of strontium (SrSO_4) falls. The formation of this precipitate is promoted by stirring and by setting the liquid aside for some time.

Barium is precipitated immediately under similar circumstances.

Third Analytical Reaction.—To a dilute solution of a strontium salt add yellow chromate of potassium; no precipitate falls.

Barium may be separated from strontium by chromate of potassium, that reagent at once precipitating barium from aqueous or acetic solutions.

Fourth Analytical Reaction.—Insert a fragment of a strontium salt in the blowpipe-flame, or other equally colourless

flame, or hold the end of a platinum wire dipped into a strontium solution in the flame; a crimson colour is imparted.

Other Analytical Reactions.—Alkali-metal phosphates, arseniates and oxalates give white insoluble precipitates with strontium as with barium and calcium.—Strontium, like calcium, but unlike barium, is not precipitated by hydrofluosilicic acid.

CERIUM. Ce. At. wt. 92.—This element occurs in the mineral cerite (a silicate of iron, calcium, and the three rare metals cerium, lanthanum, and didymium); also occasionally as impure fluoride, carbonate, and phosphate. The oxalate of cerium, a white granular powder, is the only official salt; it may be obtained from cerite by boiling the powdered mineral in strong hydrochloric acid for several hours, evaporating, diluting, and filtering to separate silica; adding ammonia to precipitate hydrates of all the metals except calcium; filtering off, washing, redissolving in hydrochloric acid, and adding oxalic acid to precipitate oxalate of cerium. The preparation will still contain oxalates of lanthanum and didymium: it is therefore strongly calcined, the resulting oxides of lanthanum and didymium dissolved out to some extent by boiling with a concentrated solution of chloride of ammonium, the residual oxide of cerium dissolved in boiling hydrochloric acid, and oxalate of ammonium added to precipitate oxalate of cerium ($\text{Ce}^{\text{C}}\text{C}_2\text{O}_4, 3\text{H}_2\text{O}$).

Oxalate of cerium (*Cerii Oxalas*, B.P.) is decomposed at a dull red heat, 48 per cent. of a yellow or, more generally, salmon-coloured mixture of oxides remaining. Usually the didymium present gives the ignited residue a reddish or reddish-brown colour. The oxides are soluble in boiling hydrochloric acid (without effervescence, indicating, indirectly, absence of earthy and other carbonates or oxalates); and the solution gives, with excess of a saturated solution of sulphate of potassium, a crystalline precipitate of double sulphate of cerium and potassium. Alumina mixed with oxalate of cerium may be detected by boiling with solution of potash, filtering, and adding excess of solution of chloride of ammonium, when a white flocculent precipitate of hydrate of aluminium will be obtained. The oxalic radical is recognised by neutralizing the potash solution by acetic acid and adding chloride of

calcium; white oxalate of calcium is then precipitated: this precipitate, though insoluble in acetic, should be wholly dissolved by hydrochloric acid.

According to H. G. Greenish most samples of oxalate of cerium contain as impurities traces of lead, iron and magnesium.

MANGANESE.

Symbol Mn. Atomic weight 55.

Source.—Manganese is a constituent of many minerals, and, as black oxide (MnO_2) (*Manganesii Oxidum Nigrum*, B. P.), or *pyrolusite* (from *πῦρ*, *pur*, fire, and *λύσις*, *lusis*, a loosing or resolving, in allusion to the readiness with which it is split up by heat into a lower oxide and oxygen), occurs frequently in abundance in the south-west of England, Aberdeenshire, and most of the countries of Europe. It is met with as a steel-grey mass of prismatic crystals, or in black shapeless lumps.

The chemical position of manganese is close to iron and three other metals still to be considered—cobalt, nickel, and chromium. Its atom apparently has sexivalent affinities, as seen in manganate of potassium (K_2MnO_4); but commonly it is quadrivalent (Mn^{IV}) or bivalent (Mn^{II}).

Uses.—Metallic manganese is only used in alloy with iron in the manufacture of some varieties of steel. The black oxide is an important agent in the production of chlorine, and in the preparation of green and red disinfecting manganates, purple glass, and black glazes for earthenware.

Reactions having both Synthetical and Analytical Interest.

First Reaction.—Boil a few grains of black oxide of manganese with some drops of hydrochloric acid until chlorine ceases to be evolved; add water, and filter; the filtrate is a solution of manganous chloride (MnCl_2).



This is the reaction commonly applied in the preparation of chlorine gas. It is also a ready method of preparing a manganous salt for analytical experiments. Coupled with the application of reagents to the filtrate, the reaction is that by which a black powder or mineral would be recognised as

black oxide of manganese. Black oxide of manganese dissolves in cold hydrochloric acid, forming a dark brown solution of a higher chloride or chlorides, MnCl_2 , Mn_2Cl_7 , or, possibly, MnCl_4 .

Second Reaction.—Heat a particle of a manganese compound with a grain or two of carbonate and hydrate of potassium and a fragment of nitrate or chlorate of potassium on platinum foil in the blowpipe-flame; a green mass containing *manganate of potassium* (K_2MnO_4) results. Boil the foil in a little water; the green manganate dissolves and soon changes to solution of the purple *permanganate of potassium* ($\text{K}_2\text{Mn}_2\text{O}_8$).

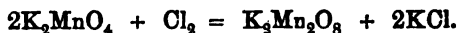
Carefully performed this is a delicate test for manganese.

The reaction is similar to that by which permanganate of potassium (*Potassæ Permanganas*, B. P.) is directed to be prepared for use in volumetric analysis. *Liquor Potassæ Permanganatis*, B. P., is a solution of 80 grains of permanganate of potassium in 1 pint of distilled water. Equations showing the exact action which occurs in making the salt according to the process of the British Pharmacopœia have already been given in connection with the compounds of potassium (*vide* p. 78). The proportions of ingredients and details of the operation are as follows:—

Reduce $3\frac{1}{2}$ parts (for experiment, each "part" may be $\frac{1}{2}$ ounce) of chlorate of potassium to fine powder, and mix it with 4 of black oxide of manganese; put the mixture into a porcelain basin, and add to it 5 parts of solid caustic potash, previously dissolved in 4 parts of water. Evaporate to dryness, stirring diligently to prevent spurting. Pulverize the mass, put it into a covered Hessian or Cornish crucible, and expose it to a dull red heat (not higher) for an hour (20 or 30 minutes for quantities of 1 or 2 ounces), or till it has assumed the condition of a semi-fused mass. Allow to cool, pulverize, and boil with about 30 parts of water. Let the insoluble matter subside, decant the fluid, boil again with about 10 parts of water, again decant, neutralize the united liquors accurately with diluted sulphuric acid (or better, carbonic acid gas), and evaporate till a pellicle forms. Set aside to cool and crystallize. Drain the crystalline mass, boil it in 6 parts of water, and strain through a funnel the throat of which is lightly obstructed by a little asbestos or

gun-cotton. Let the fluid cool and crystallize, drain the dark purple slender prismatic crystals, and dry them by placing under a bell jar over a vessel containing sulphuric acid.

Instead of converting the manganate into permanganate by ebullition, and neutralizing the free alkali by acid, by which one-third of the manganese is lost, Städeler recommends chlorine to be passed through the cold solution until the green colour is entirely changed to purple.



Solutions of the manganates of potassium and sodium are in common use as disinfectants under the name of Condyl's fluid. They act by oxidizing organic matter, the manganic or permanganic radical being reduced to black manganic oxide, or even a lower oxide. The reason for using asbestos instead of paper in filtering the solutions will now be understood.

The changes in colour which the green mass of the above process undergoes when dropped into warm water produced for it the old name of *mineral chameleon*.

Third Reaction.—Make a borax bead by heating a fragment of the salt on the looped end of a platinum wire in the blow-pipe-flame until a clear transparent globule is obtained. Place on the bead a minute portion of a manganese compound, or touch it with a drop of solution. Again fuse the borax; a bead of a violet or amethystine tint is produced.

This is a good analytical reaction. It has also synthetical interest, illustrating the use of black oxide of manganese in producing common purple-tinted glass.

Expose the bead to the reducing part of the flame (p. 146), the part nearer to the blowpipe, where there are highly heated hydrocarbon gases greedy of oxygen; the colour disappears.

This is owing to the reduction of the manganic compound to a manganous condition, in which it no longer possesses peculiar colouring-power. This action also illustrates the use of black oxide of manganese in glass-manufacture. Glass when first made is usually of a green tint, owing to the presence of ferrous impurities; the addition of manganic oxide to the materials converts the ferrous into ferric compounds, which have comparatively little colorific power, it itself being

thereby reduced to manganous oxide, which also gives but little colour. If excess of manganic oxide be added, a purple tint is produced.

Reactions having Analytical Interest (Tests).

Fourth Reaction.—Through a solution of a manganous salt acidified by hydrochloric acid pass sulphuretted hydrogen; no decomposition occurs. Add ammonia; the sulphhydrate of ammonium thus formed causes the precipitation of a yellowish-pink or flesh-tinted precipitate of manganous sulphide (MnS) in a hydrous state.

This reaction is characteristic, sulphide of manganese being the only flesh-coloured sulphide known. The salt used may be the manganous chloride obtained in the first reaction; but such crude solutions usually give a black precipitate with sulphhydrate of ammonium, owing to the presence of iron. The latter element may be precipitated, however, on adding excess of ammonia (and rapidly filtering, or oxygen will be absorbed and most of the manganese also be precipitated) or on boiling the manganous solution with a very little carbonate of sodium, which attacks the ferric salt in preference to the manganous. Pure manganous chloride may be similarly obtained on boiling the impure solution with manganous carbonate; the latter decomposes the ferric chloride with production of ferric hydrate and more manganous chloride, and evolution of carbonic acid gas.

To the recently precipitated manganous sulphide add acetic acid; it is dissolved.

This solubility enables manganese to be separated from nickel, cobalt, and zinc, whose sulphides are insoluble in weak acetic acid. To express the fact in another way—manganese is not precipitated by sulphuretted hydrogen from a solution containing free acetic acid only.

Fifth Reaction.—To solution of a manganous salt add ammonia; a white precipitate of manganous hydrate (Mn_2HO) falls. Add excess of ammonia; some of the precipitate is dissolved and may be detected in the quickly filtered solution by sulphhydrate of ammonium. But both precipitate and

solution rapidly absorb oxygen, the manganese passing into a more highly oxidized condition in which it is insoluble in ammonia.

The fixed alkalies give a similar precipitate *insoluble* in excess. The precipitate rapidly absorbs oxygen, becomes brown, and gradually passes into a higher oxide.

Many other reactions occur between manganese salts and various reagents, but are of no particular synthetical or analytical interest.

A good method, proposed by Crum, for detecting minute quantities of manganese consists in adding diluted nitric acid, and either red lead or the puce-coloured oxide or peroxide of lead, to the solution, and then boiling; a red tint, said to be due to permanganic acid, is imparted to the liquid. If chlorides are present the manganese, etc., should be separated by potash or soda, the precipitate be well washed, dissolved in nitric acid, and then the oxide be added.

COBALT.

Symbol Co. Atomic weight 58.8.

Source.—Cobalt occurs sparingly in nature as the arsenide (CoAs_3) or *tin-white cobalt*, and occasionally as a double arsenide and sulphide (CoAs_3 , CoS_2), or *cobalt-glance* (from *glanz*, brightness, in allusion to its lustre).

Uses.—Its chief use is in the manufacture of blue glass, the colour of which is due to a compound of cobalt. Cobalt is also the colouring constituent of *smalt* (from *smelt*, a corruption of *melt*), a finely ground sort of glass used as a blue pigment by paper-stainers and others, and employed also by laundresses to neutralize the yellowish appearance of washed linen.

The salts of cobalt may be obtained from the oxide (CoO), and the oxide from *saffre*, a mixture of sand and roasted ore.

Quantivalence.—The atom of cobalt often exhibits quadrivalent affinities, but still more often exerts only bivalent powers (Co^{II}). Cobalt has analytical relations with zinc, nickel, and manganese, and may be regarded as a member of the iron group.

Analytical Reactions (Tests).

First Analytical Reaction.—Pass sulphuretted hydrogen through a solution of a salt of cobalt—the chloride (CoCl_2) or nitrate (Co_2NO_3) for example; no decomposition occurs. Add ammonia; the sulphhydrate of ammonium thus formed causes the precipitation of black sulphide of cobalt (CoS).

The moist precipitate slowly absorbs oxygen from the air, yielding some sulphate of cobalt (CoSO_4).

Second Analytical Reaction.—Add ammonia gradually to a cobalt solution; a blue precipitate of impure hydrate of cobalt (Co_2HO) falls. Add excess of ammonia; the precipitate is dissolved, yielding a liquid somewhat more reddish brown than the original solution.

A similar precipitate is given by the fixed alkalies, *insoluble* in excess.

Third Analytical Reaction.—Make a borax bead by heating a fragment of the salt on the looped end of a platinum wire in the blowpipe-flame until a clear transparent globule is obtained. Place on the bead a minute portion of cobalt compound, or touch it with a drop of solution. Again fuse the borax; a blue bead results.

This is a delicate test for cobalt. From what has previously been said, it will be seen that this experiment has also considerable synthetical interest.

Fourth Analytical Reaction.—To a solution of a salt of cobalt add two or three drops of hydrochloric acid, then excess of solution of cyanide of potassium, and boil for ten minutes; oxygen is absorbed, and cobalticyanide of potassium ($\text{K}_6\text{Co}_3\text{Cy}_{12}$) formed. Add hydrochloric acid, and boil the mixture (in a fume-cupboard, to avoid inhalation of any hydrocyanic acid); the excess of cyanide of potassium is thus decomposed, but the cobalticyanide unaffected. Now add *excess* of solution of potash; the cobalticyanide of potassium is decomposed, but the cobalt remains dissolved in the alkaline liquid.

Nickel under similar circumstances is precipitated, the reaction thus affording means of separating these closely allied metals from each other.

Other reactions between a cobalt solution and different reagents may be performed, and various precipitates obtained; but these have no special analytical interest.

Invisible Ink.—Many salts of cobalt containing water of crystallization are light red, the anhydrous more or less blue. Prove this by writing some words on paper with a solution of chloride of cobalt sufficiently dilute for the characters to be invisible when dry: hold the sheet before a fire or over a flame; the letters at once become visible, distinct, and of a blue colour. Breathe on the words, or set the sheet aside for a while; the characters are once more invisible, owing to absorption of moisture. Hence solution of chloride of cobalt forms one of the so-called *sympathetic inks*.

NICKEL.

Symbol Ni. Atomic weight 58.8.

Nickel is, chemically, closely allied to cobalt, the ores of the two metals being, commonly, associated in nature. Indeed it is from *speiss*, an arsenio-sulphide of nickel obtained in the manufacture of a pigment of cobalt, smalt, already mentioned, that most of the nickel met with in commerce is obtained. It is much used in the preparation of the white alloy known as German or nickel silver.

Quantivalence.—Nickel exerts bivalent activity (Ni^{+2}) in its ordinary compounds. Its salts and their solutions are usually green. They are chiefly made, directly or indirectly, from the metal itself.

Analytical Reactions (Tests).

First Analytical Reaction.—Pass sulphuretted hydrogen through a solution of a salt of nickel—chloride (NiCl_2), nitrate (Ni_2NO_3), or sulphate (NiSO_4); no decomposition occurs. Add ammonia; the sulphhydrate of ammonium thus formed causes the precipitation of black sulphide of nickel (NiS).

Note.—When sulphide of nickel is precipitated by the direct addition of the common yellow solution of sulphhydrate of ammonium, which always contains free sulphur, there is much difficulty in filtering the mixture, owing to the slight solubility of sulphide of nickel in the reagent and the formation of some sulphate of nickel (NiSO_4), oxygen being absorbed from the air by the sulphide. This may be avoided by warming the mixture and using freshly made sulphhydrate of ammonium, in which the sulphide of nickel is insoluble; or, when practicable, the salt of nickel may be precipitated from an ammoniacal solution by sulphuretted hydrogen.

Second Analytical Reaction.—Add ammonia drop by drop to a nickel solution; a pale green precipitate of hydrate of nickel (Ni_2HO) falls—especially on boiling the mixture. Add excess of ammonia; the precipitate dissolves, yielding a bluish rather than the original green-coloured solution.

A similar precipitate is given by the fixed alkalies, *insoluble* in excess.

Third Analytical Reaction.—Nickel salts colour a borax bead, when hot, a reddish-yellow tint; the reaction is not very serviceable analytically.

Fourth Analytical Reaction.—To a solution of a salt nickel add solution of cyanide of potassium; cyanide of nickel (NiCy_2) is precipitated. Add excess of solution of cyanide of potassium; the precipitate is dissolved with formation of double cyanide of nickel and potassium ($\text{NiCy}_2, 2\text{KCy}$). Next add hydrochloric acid, and boil the mixture (in a fume-cupboard), adding a little hydrochloric acid from time to time until all smell of hydrocyanic acid has disappeared. Lastly add *excess* of solution of potash; hydrate of nickel is precipitated.

Qualitative Separation of Cobalt and Nickel.

The foregoing reaction serves for the separation of nickel from cobalt. On adding excess of hydrochloric acid to a solution containing the two metals, together with cyanide of potassium, a precipitate of cyanide of nickel and cobalticyanide of nickel occurs. By ebullition with excess of hydro-

chloric acid the cyanide of nickel is decomposed, chloride of nickel going into solution. On then adding excess of potash, hydrate of nickel is precipitated. The cobalticyanide of nickel is not decomposed by the acid; but it is by the alkali, its cobalt going into solution, and its nickel remaining insoluble as hydrate. After filtering off the nickel, cobalt is detected in the filtrate by evaporating to dryness and testing the residue with borax in the blowpipe-flame.

W. R. Dunstan thus modifies this process:—

To the acidified solution containing the two metals, potassium cyanide is added in excess until a clear solution is obtained. To a portion of this liquid a few drops of yellow ammonium sulphide are added; a dark red coloration indicates cobalt. This reaction is very characteristic, and is not interfered with by the presence of nickel. The solution does not give the reaction after exposure to the air or after ebullition. To another portion of the liquid some sodium hypobromite solution is added, and the solution gently warmed; a black precipitate indicates nickel. The remainder of the above solution is boiled for a few minutes, and when cool hydrochloric acid added in excess; if both metals are present a dense bluish-white precipitate occurs. The solution is boiled for some time till free from hydrocyanic acid, the precipitate remaining insoluble, which is a sure indication of the presence of both metals. To this liquid containing the precipitate, large excess of potassium hydrate is added. The precipitate of nickelous cobalticyanide disappears, and is replaced by a pale green flocculent precipitate of nickelous hydrate.

The reducing action of glycerine on nickelic hydrate, $\text{Ni}_2\text{6HO}$, and the absence of any similar action on cobaltic hydrate affords another method of separating the metals. (*See Pharmaceutical Journal*, May 31, 1879.)

Another process.—To any solution containing cobalt and nickel add excess of ammonia, and then enough solution of ferridecyanide of potassium to dissolve any precipitate. If cobalt be present a reddish-brown colour is produced in the liquid, due to the formation and solution of ferridecyanide of cobalt. Now add solution of ferrocyanide of potassium; a whitish precipitate of ferrocyanide of nickel appears, either at once, or, if the liquid is very weak, on *partially* neutralizing the free ammonia by hydrochloric acid. The ammonia

must not be entirely neutralized by acid, nor enough acid added to decompose the reddish ferridcyanide of cobalt. If these precautions be not observed, a greenish-white precipitate of ferrocyanide of cobalt may be formed, and be mistaken for the ferrocyanide of nickel.

The value of this method (Skey and Davies) turns on the facts that ferridcyanide of nickel is not a coloured body, while ferridcyanide of cobalt is reddish brown, and that ferridcyanogen has, apparently, in *ammoniacal solution*, greater affinity for cobalt than for nickel, while ferrocyanogen has, apparently, greater affinity for nickel than for cobalt. The formulæ of these so-called ferrocyanides and ferridcyanides of cobalt and nickel have not been definitely ascertained.

Other reactions. between a nickel solution and various reagents give, in many cases, insoluble precipitates which, from their green colour, are occasionally useful in distinguishing nickel from allied elements.

CHROMIUM.

Symbol Cr. Atomic weight 52.5.

Source.—The chief ore of chromium is chrome ironstone (a mixture of the metallic oxides FeO , Cr_2O_3), occurring chiefly in the United States and Sweden. In constitution it seems to resemble magnetic iron ore (FeO , Fe_3O_4).

Preparation of Red Chromate of Potassium.—On roasting the powdered ore with carbonate of potassium and nitre, yellow chromate of potassium (K_2CrO_4) is obtained; the mass, treated with acid, yields red or bichromate ($\text{K}_2\text{Cr}_2\text{O}_7$, Cr_2O_3) (*Potassa Bichromas*, B.P.); from this salt other chromates are prepared, and by reduction, as presently explained, the salts of chromium itself. The yellow and orange chromates of lead are largely used as pigments.

Notes.—Red chromate of potassium is a somewhat abnormal salt, containing, probably, neutral chromate associated with chromic anhydride. The value of chromates as chemical reagents is alluded to in connection with chromate of barium (pp. 111 and 124). Heated strongly in a crucible, red chromate of potassium splits up into yellow chromate, glistening oxide of chromium, and oxygen.

Quantivalence.—Chromium stands in close chemical relation to iron, aluminium, and manganese. Its atom is sexivalent if the formula of the fluoride (CrF_6) be correct. Like iron and aluminium, it is trivalent, as seen in chromic chloride (Cr_2Cl_6), but sometimes exerts only bivalent activity, as in chromous chloride (CrCl_2).

Passage of chromium from the acidulous to the basylous side of salts. Through an acidified solution of red chromate of potassium pass sulphuretted hydrogen; sulphur is deposited, and a green salt of chromium remains in solution—chloride (Cr_2Cl_6) if hydrochloric acid be used, and sulphate (Cr_2SO_4) if sulphuric be the acid employed. Boil the liquid to expel excess of sulphuretted hydrogen, filter, and reserve the solution for subsequent experiments. (For an equation explanatory of this reaction, see p. 273.)

Alcohol, sugar, or almost any substance which is tolerably liable to oxidation will answer as well as sulphuretted hydrogen.

Sulphate of chromium (Cr_2SO_4), like sulphate of aluminium (Al_2SO_4), unites with alkali-metal sulphates to form *alums*, which resemble common alum both in crystalline form and, as far as we know, in internal structure: they are of a purple colour.

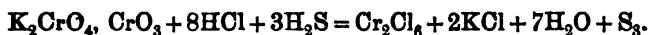
REACTIONS.

Chromium as chromic acid or other chromate. This is the state in which chromium will usually be met with, the most common salt being the red chromate or bichromate of potassium. Mix four volumes of a cold saturated aqueous solution of red chromate of potassium with five of oil of vitriol; on cooling, *chromic anhydride* (CrO_3) (*Acidum Chromicum*, U. S. P.) separates in crimson needles. After well draining, the crystals may be freed from adhering sulphuric acid by washing once or twice with nitric acid: the latter may be removed by passing dried and slightly warmed air through a tube containing the crystals. In contact with moisture chromic anhydride takes up water and forms solution of true

chromic acid (H_2CrO_4). Chromic anhydride is a powerfully corrosive oxidizing agent; it melts between 356° F. and 374° F.

The oxygen in chromic acid and other chromates, and in manganates, permanganates, black oxide of manganese, and puce-coloured oxide of lead, is in a physically different state to that in peroxide of hydrogen, peroxide of barium, and similar compounds. On bringing chromic acid, or the above acidified solution of red chromate of potassium, into contact with solution of peroxide of hydrogen a strong effervescence of oxygen ensues. According to Schönbein and Brodie the oxygen in chromic acid is in the negative or ozonic state, while that of peroxide of hydrogen is in the positive or so-called antozonic condition. Both are equally active, but neutralize each other, forming neutral or ordinary oxygen.

In the analytical examination of solutions containing chromates, the chromium will always come out in the state of green chromic hydrate along with ferric hydrate and alumina, the prior treatment by sulphuretted hydrogen reducing the chromium in the molecule to the lower state, thus:—



Chromium having been found in a solution, its condition as chromate may be ascertained by applying to the original solution salts of barium, mercury, lead, and silver. (See the various paragraphs relating to those metals.)

Ba_2NO_3 gives yellow $BaCrO_4$ with chromates.

$HgNO_3$ " red Hg_2CrO_4 " "

$AgNO_3$ " red Ag_3CrO_4 " "

" " " Ag_3CrO_4, CrO_3 with bichromates.

$Pb_2C_2H_3O_2$ " yellow $PbCrO_4$ with both.

Nitrate of barium does not completely precipitate bichromates, bichromate of barium being soluble in water; the chromate of barium is insoluble in water or acetic acid, but soluble in hydrochloric or nitric acid. Mercurous nitrate does not wholly precipitate bichromates: mercuric nitrate or chloride only partially precipitates chromates, and does not precipitate bichromates. The mercurous chromate is insol-

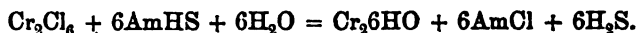
uble, or nearly so, in diluted nitric acid. The silver chromates are soluble in acids and alkalies. Acetate of lead precipitates chromates and bichromates, acetic acid being set free in the latter case.

A delicate reaction for dry chromates will be found in the formation of *chlorochromic anhydride* (CrO_2Cl_2). A small portion of the chromate is placed in a test-tube with a fragment of dry chloride of sodium and a drop or two of oil of vitriol, and the mixture heated; red irritating fumes of chlorochromic anhydride are evolved and condense in dark red drops on the side of the tube.

Larger quantities of pure distilled chlorochromic anhydride are obtained by the same reaction, the operation being conducted in a retort, with thoroughly dry materials, for the compound is decomposed by water. It may be regarded as chromic anhydride in which an atom of oxygen is displaced by an equivalent quantity (two atoms) of chlorine. It is not used in medicine, but is of interest to the chemical student as being an illustration of a large class of similar bodies—*chloro-acidulous* or *chloro-anhydro compounds*. The reaction is also occasionally serviceable for the detection of chlorides.

Analytical Reactions of Chromium Salts (Tests).

First Analytical Reaction.—To solution of a salt of chromium (chloride, sulphate, or chrome alum) add sulphhydrate of ammonium; a bulky green precipitate of chromic hydrate (Cr_26HO), containing a large quantity of water (7 molecules, $7\text{H}_2\text{O}$), is precipitated.



Second Analytical Reaction.—To solution of a chromium salt add ammonia; chromic hydrate is precipitated, insoluble in excess.

Third Analytical Reaction.—To solution of a chromium salt add solution of potash or soda drop by drop; chromic hydrate is precipitated. Add excess of the fixed alkali: the precipitate is dissolved. Well boil the solution; the chromic hydrate is reprecipitated.

Iron, Chromium, and Aluminium Salts, chemically so alike, may be separated by this reaction. Ferric hydrate is insoluble in solutions of the fixed alkalies, cold or hot; chromium hydrate soluble in cold but not in hot: hydrate of aluminium in both. To a solution containing all three metals, therefore, add potash or soda, stir, and filter; the iron is thrown out, boil the filtrate, and filter; the chromium is thrown out: neutralize the filtrate by acid, and then add ammonia; the aluminium is thrown out. Note, however, that ferric hydrate will prevent hydrate of chromium being dissolved by potash or soda if the ferric hydrate is in considerable excess. Before concluding that chromium is entirely absent the fourth reaction should be performed. The hydrates of iron, chromium, and aluminium are insoluble in ammonia, and may therefore be easily separated from the hydrates of the somewhat analogous metals zinc, cobalt, nickel, and manganese.

Fourth Analytical Reaction.—Add a salt of chromium (either of the above precipitates of chromic oxide or the dry residue of the evaporation of a few drops of a solution of a chromium salt) to a few grains of nitre and carbonate of sodium on platinum foil, and fuse the mixture in the blowpipe-flame; a yellow mass of chromate of potassium and sodium (KNaCrO_4) is formed. Dissolve the mass in water, add acetic acid to decompose excess of carbonate, and apply the reagents for chromates.

This is a delicate and useful reaction if carefully performed.

TIN.

Symbol Sn. Atomic weight 118.

Source.—The chief ore of tin is stannic oxide (SnO_2), occurring in veins under the name of *tinstone*, or in alluvial deposits as *stream-tin*. The oldest mines are those of Cornwall. Much tin is now imported from Australia.

Preparation.—The metal is obtained by reducing the roasted and washed ore by charcoal or anthracite* coal at a high tem-

* *Anthracite* (from *ἀνθραξ*, *anthraz*, a burning coal) or *stone coal* differs from the ordinary *bituminous* or *caking coal* in containing less volatile matter, and, therefore, in burning without flame. It gives a higher temperature, and from its non-caking properties is, in furnace operations, more manageable than bituminous coal.

perature, and is purified by slowly heating, when the pure tin, fusing first, is run off, a somewhat less fusible alloy of tin with small quantities of arsenicum, copper, iron, or lead remaining. The latter is known as *block tin*; the former heated till brittle and then hammered or let fall from a height splits into prismatic fragments resembling starch or columnar basalt, and is named *dropped* or *grain tin*. Good tin emits a crackling noise in bending, termed the cry of tin, caused by the friction of its crystalline particles on each other.

Uses.—Tin is an important constituent of such alloys as pewter, Britannia metal, solder, speculum-metal, bell-metal, gun-metal, and bronze. It is very ductile, and may be rolled into plates or leaves, known as *tin foil*, varying from $\frac{1}{10}$ to $\frac{1}{1000}$ of an inch in thickness. Common tin foil, however, usually contains a large proportion of lead. The reflecting surface of *looking-glasses* was, formerly, always an amalgam of tin and mercury, produced by carefully sliding a plate of glass over a sheet of tin foil on which mercury had been rubbed and then excess of mercury poured—but pure silver, deposited from a solution, is now largely employed. *Pins* are made of brass wire on which tin is deposited. *Tin plate*, of which common utensils are made, is iron alloyed with tin by dipping the acid-cleansed sheet into melted tin covered with oil, which, by dissolving any trace of oxide, or, perhaps, by preventing oxidation, enables the tin more completely to alloy with the iron. *Tin tacks* are in reality tinned iron tacks, a tin nail would be too soft to drive into wood. Tin may be granulated by melting and triturating briskly in a hot mortar, by shaking melted tin in a box on the inner sides of which chalk has been rubbed, or, in thin little bells or corrugated fragments (Granulated Tin, B. P.), by melting in a ladle and, as soon as fluid, pouring from the height of a few feet into water. Powdered tin has been used medicinally as a mechanical irritant to promote expulsion of worms. The hairs of the pod of *Kiwach* or *Cowhage* (Hindustani) (*Mucuna pruriens*) (P. I.) is almost the only other medicine (excluding diluents and dentifrices) which acts in such a directly mechanical manner.

The chemical position of tin among the metals is close to that of arsenicum and antimony. Its atom is quadrivalent and bivalent. The two classes of salts are termed stannic and stannous respectively. They are all made directly or indirectly from the metal itself.

REACTIONS HAVING (a) SYNTHETICAL AND (b) ANALYTICAL
INTEREST.

(a) *Synthetical Reactions.*

Chloride of Tin. Stannous Chloride.

First Synthetical Reaction.—Warm a fragment of tin with hydrochloric acid; hydrogen escapes and solution of stannous chloride (SnCl_2) is formed. It may be retained for future experiments.

One ounce of tin dissolved in three fluid ounces of hydrochloric acid and one of water, and the resulting solution diluted to five fluid ounces, constitutes the "Solution of Chloride of Tin," B. P.

Solid stannous chloride.—By evaporation of the above solution stannous chloride is obtainable in crystals ($\text{SnCl}_2 \cdot 2\text{H}_2\text{O}$). It is a powerful reducing agent, even a dilute solution precipitating gold, silver, and mercury from their solutions, converting ferric and cupric into ferrous and cuprous salts, and partially deoxidizing arsenic, manganic, and chromic acids. It absorbs oxygen from the air, and is decomposed when added to a large quantity of water unless some acid be present. It is used as a mordant in dyeing and calico-printing.

Perchloride of Tin. Stannic Chloride.

Second Synthetical Reaction.—Through a portion of the solution of the stannous chloride of the previous reaction pass chlorine gas; solution of stannic chloride (SnCl_4), is formed. Or add hydrochloric acid to the stannous solution, boil, and slowly drop in nitric acid until no more fumes are evolved; again stannic chloride results. Reserve the solutions for subsequent experiments.

Stannic Oxide or Anhydride and Stannates.

Third Synthetical Reaction.—Boil a fragment of tin with nitric acid, evaporate to dryness, and strongly calcine the residue; light buff-tinted stannic anhydride (SnO_2) is produced. Heat the stannic anhydride with excess of solid caustic potash or soda; stannate of the alkali metal (K_2SnO_3)

or Na_2SnO_3) results. Dissolve the stannate in water, and add hydrochloric acid; white, gelatinous *stannic acid* (H_2SnO_3) is precipitated. Stannic acid is also obtained on adding an alkali to solution of stannic chloride: it is soluble in excess of acid or alkali.

The product of the action of nitric acid on tin is also an acid, but, from its insolubility in hydrochloric and other acids, is different from ordinary stannic acid. It is termed *meta-stannic acid* (from *meta*, beyond), and probably has a composition expressed by the formula $\text{H}_{10}\text{Sn}_5\text{O}_{15}$ (*vide* Index, "Isomerism"). It is also produced on gently heating stannic acid:—



Metastannates have the general formula $\text{M}_2\text{H}_3\text{Sn}_5\text{O}_{15}$.

Both acids yield buff-coloured stannic oxide or anhydride (SnO_2) when strongly heated. The latter is employed in polishing plate under the name of *putty powder*. *Stannate of sodium* ($\text{Na}_2\text{SnO}_3 \cdot 4\text{H}_2\text{O}$) is used as a mordant by dyers and calico-printers under the name of *tin prepare-liquor*.

(b) Reactions having Analytical Interest (Tests).

Stannous or stannic salts.—Heat any solid tin compound with a mixture of cyanide of potassium and carbonate of sodium on charcoal by the inner flame of the blowpipe. Globules of tin separate, having, when cut by a knife, characteristic brightness and hardness.

STANNOUS SALTS.

First Analytical Reaction.—Through a diluted solution of a stannous salt (stannous chloride, for example) pass sulphuretted hydrogen gas; brown stannous sulphide (SnS) is precipitated. Pour off the supernatant liquid, add ammonia to the moist precipitate (to neutralize acid), and, lastly, yellow sulphhydrate of ammonium solution; the precipitate is dissolved.

Aqueous solution of sulphhydrate of ammonium becomes yellow when a day or two old, and then contains excess of sul-

phur, that element having become displaced by oxygen absorbed from the air: hence in the above reaction the stannous sulphide (SnS), in dissolving, becomes stannic sulphide (SnS_2); for the latter is precipitated on decomposing the alkaline liquid by an acid.

Second Analytical Reaction.—To solution of a stannous salt add solution of potash or soda; white stannous hydrate falls (Sn_2HO). Add excess of the alkali; the precipitate dissolves. Boil the solution; some of the tin is reprecipitated as blackish stannous oxide (SnO).

Ammonia gives a similar white precipitate, insoluble in excess. The alkaline carbonates do the same, carbonic acid gas escaping.

STANNIC SALTS.

Third Analytical Reaction.—Through solution of a stannic salt (stannic chloride, for example) pass sulphuretted hydrogen gas; yellow stannic sulphide (SnS_2) is precipitated. Pour off the supernatant liquid, and to the moist precipitate add ammonia (to neutralize acid), and then sulphhydrate of ammonium; the precipitate dissolves.

Note.—In precipitating stannic sulphide the presence of too much hydrochloric acid must be avoided; the formation of the precipitate is also facilitated if the solution be warmed. Stannic sulphide, like the sulphides of arsenicum and antimony, dissolves in a solution of alkaline sulphide or sulphhydrate, with formation of definite crystallizable *sulphostannates* (M_2SnS_3).

Anhydrous stannic sulphide, prepared by sublimation, has a yellow or orange lustrous appearance, and is used by decorators as *bronzing-powder*. It is sometimes termed *mosaic gold*.

Fourth Analytical Reaction.—To solution of a stannic salt add potash or soda; white stannic acid falls (H_2SnO_3). Add excess of the alkali; the precipitate dissolves. Boil the mixture; no reprecipitation occurs—a fact enabling stannic to be distinguished from stannous salts.

Ammonia gives a similar precipitate, soluble, but not readily, in excess. The fixed alkali-metal carbonates do the same,

carbonic acid gas escaping; after a time the stannic salt is again deposited, probably as stannate of the alkali-metal. Carbonate of ammonium and acid carbonates of alkali-metals give a precipitate of stannic acid insoluble in excess.

Antidotes.—In cases of poisoning by tin salts (dyers' tin-liquor *e.g.*), solution of carbonate of ammonium should be given. White of egg is also said to form an insoluble precipitate with compounds of tin. Vomiting should be speedily induced, and the stomach-pump quickly applied.

GOLD.

Symbol Au. Atomic weight 196.7.

Source.—Gold occurs in the free state in nature, occasionally in nodules or *nuggets*, but commonly in a finer state of division termed *gold dust*.

Preparation.—Gold is separated from the sand, crushed quartz, or other earthy matter with which it may be associated by agitation with water, when the gold, from its relatively greater specific gravity, falls to the bottom of the vessel first, the lighter mineral matter being allowed to run off with the water. From this rich sand the gold is dissolved out by mercury, the amalgam filtered and afterwards distilled, when the mercury volatilizes and gold remains. The amalgamation may be much facilitated by the use of a small proportion of sodium, as already described in treating of silver.

Pure gold is too soft for general use as a circulating medium. *Gold coin* is an alloy of copper and gold, that of Great Britain containing 1 of the former to 11 of the latter, or $8\frac{1}{3}$ per cent. of copper, that of France, Germany, and the United States about 10 per cent. *Jewellers' gold* varies in quality, every 24 parts containing 18, 15, 12, or 9 parts of gold, the alloys being technically termed 18, 15, 12, or 9 *carat fine*. Articles made of the better qualities are usually stamped by authority. Trinkets of inferior intrinsic worth are commonly thinly coated with pure gold by electro-deposition or otherwise. *Gold leaf* is nearly pure gold passed between rollers till it is about $\frac{1}{100}$ of an inch in thickness and then hammered between sheets of animal membrane, termed gold-beaters' skin and calf-skin vellum, till it is $\frac{1}{100000}$ or $\frac{1}{100000}$ of an inch in thickness. It may even be hammered till 280,000 leaves would be required to form a pile an inch thick.

Gold Coinage.—The weight of gold is expressed in this country in ounces troy and decimal parts of an ounce; and the metal is always taken to be of standard fineness (11 gold and 1 alloy), unless otherwise described. The degree of fineness of gold, as ascertained by assay, is expressed decimally, fine pure gold ("gold free from metallic impurities," B. P.) being taken as unity, or 1·000. Thus gold of British standard is said to be 0·9166 fine, of French standard 0·900 fine. The legal weight of the sovereign is 0·2568 ounce of standard gold, or 123·274 grains. The weight came from one pound of standard gold (5760 grains) being coined into $44\frac{1}{4}$ guineas. Gold coins are legal tender to any amount, provided that the weight of each sovereign does not fall below 122·5 grains, or in the case of a half-sovereign 61·125 grains; these are the "least current" weights of the coins.

Note.—In chemical analysis gold comes out among the sulphides of the metals precipitated by sulphuretted hydrogen; and of those sulphides, it, like the sulphides of tin, antimony, and arsenicum, is soluble in yellow sulphhydrate of ammonium solution.

Quantivalence.—Gold is trivalent (Au^{+++}), but in some compounds univalent (Au^+).

REACTIONS.

Synthetical Reaction.—Place a fragment of gold (e.g. gold leaf) in ten or twenty drops of aqua regia (a mixture of three parts of nitric and four or five of hydrochloric acid), and set the test-tube aside in a warm place; solution of perchloride of gold or auric chloride (AuCl_3) results. When the metal is dissolved, evaporate nearly to dryness to remove most of the excess of acid, dilute with water, and retain the solution for subsequent experiments. Sixty grains of gold treated thus, and the resulting chloride dissolved in five ounces of distilled water, constitutes "Solution of Chloride of Gold," B. P.



This reaction has analytical interest also; for in examining a substance suspected to be or to contain metallic gold, solution would have to be effected in the above way before reagents could be applied. Gold is insoluble in hydrochloric, nitric, and the weaker acids.

Analytical Reactions (Tests).

First Analytical Reaction.—Through a few drops of solution of an auric salt (the chloride, AuCl_3 , is the only convenient one) pass sulphuretted hydrogen; brown auric sulphide (Au_2S_3) is precipitated. Filter, wash, and add yellow sulphhydrate of ammonium solution; the precipitate dissolves.

Second Analytical Reaction.—To solution of a salt of gold add ferrous chloride or sulphate, and set the tube aside; metallic gold is precipitated, a ferric salt remaining in solution.

This is a convenient way of preparing pure gold, or *fine gold* as it is termed, or of working up the gold residues of laboratory operations. The precipitate, after boiling with hydrochloric acid, washing, and drying, may be obtained in a button by mixing with an equal weight of borax or acid sulphate of potassium and fusing in a good furnace.

Third Analytical Reaction.—Add a few drops of dilute solutions of stannous and stannic chloride to a considerable quantity of distilled water; pour the liquid, a small quantity at a time, into a very dilute solution of auric chloride (AuCl_3), well stirring; the mixture assumes a purple tint, and flocks of a precipitate, known as the *Purple of Cassius* (from the name of the discoverer, M. Cassius), are produced.

The same compound is formed on immersing a piece of tin foil in solution of auric chloride; it is said to be a mixture of auric, aurous, stannic, and stannous oxides. It is the colouring agent in the finer varieties of ruby glass.

PLATINUM.

Symbol Pt. Atomic weight 198.

Source.—Platinum, like gold, usually occurs in nature in the free state, the chief sources of supply being Mexico, Brazil, and Siberia. It is separated from the alluvial soil by washing.

Uses.—The chief use of platinum is in the construction of foil, wire, crucibles, spatulas, capsules, evaporating-dishes, and stills, for the use of the chemical analyst or manufacturer.

It is tolerably hard, fusible with very great difficulty, not dissolved by hydrochloric, nitric, or sulphuric acid, and only slightly affected by alkaline substances. It is attacked by aqua regia with production of perchloride of platinum or platinic chloride (PtCl_4). It forms fusible alloys with lead and other metals, and with phosphorus a phosphide which easily melts. Neither of these substances, therefore, nor mixtures which may yield them, should be heated in platinum vessels.

*The chemical position of platinum among the elements is close to that of gold. Its atom is quadrivalent in some compounds, in others apparently bivalent (Pt''). The higher salts are termed *platinic*, the lower *platinous*.*

The specific gravity of platinum is 21.5; and that of iridium, an allied metal, 22.4.

REACTIONS.

Perchloride of Platinum. Platinic chloride.

Synthetical Reaction.—Place a fragment of platinum in a little aqua regia and set the vessel aside in a warm place, adding more acid from time to time if necessary; solution of perchloride of platinum (PtCl_4) results. Evaporate the solution to remove excess of acid, and complete the desiccation over a water-bath. Dissolve the residue in water, and retain the solution for subsequent experiments, and as a reagent for the precipitation of salts of potassium and ammonium.

A quarter of an ounce of platinum treated in the above manner, and the resulting chloride dissolved in five ounces of water, constitutes "Solution of Perchloride of Platinum," B. P.

This reaction has analytical interest also; for in examining a substance suspected to be or to contain metallic platinum, solution would have to be thus effected before reagents could be applied.

Analytical Reactions (Tests).

First Analytical Reaction.—Through a few drops of a solution of a platinic salt (PtCl_4 is the only convenient one) to which an equal quantity of solution of chloride of sodium has been added, pass sulphuretted hydrogen; dark brown platinic

sulphide (PtS_2) is precipitated. Filter, wash, and add sulphate of ammonium; the precipitate dissolves.

If chloride of sodium be not present in the above reaction, the precipitated sulphide will contain platinous chloride, and may detonate if heated.

Second Analytical Reaction.—Add excess of solution of carbonate of sodium and some sugar to solution of perchloride of platinum and boil; a precipitate of metallic platinum falls.

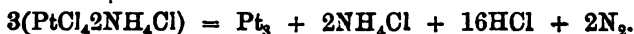
Platinum black (B. P.) is the name of this precipitate. It possesses in a high degree a quality common to many substances, but largely possessed by platinum, namely that of absorbing or occluding gases. In its ordinary state, after well washing and drying, it absorbs from the air and retains many times its bulk of oxygen. A drop of ether or alcohol placed on it is rapidly oxidized, the platinum becoming hot. This action may be prettily shown by pouring a few drops of ether into a beaker (one having portions of the top and sides broken off answers best), loosely covering the vessel with a card, and suspending within the beaker a platinum wire, one end being attached to the card by passing through its centre, the other terminating in a short coil or helix near the surface of the ether: on now warming the helix in a flame and then rapidly introducing it into the beaker, it will become red-hot and continue to glow so long as there is ether in the vessel. In this experiment real combustion goes on between the ether vapour and the concentrated oxygen of the air, the products of the oxidation revealing themselves by their odour.

Third Analytical Reaction.—To solution of perchloride of platinum add solution of chloride of ammonium; a yellow granular precipitate of double chloride of platinum and ammonium ($\text{PtCl}_4, 2\text{AmCl}$) falls. When slowly formed in dilute solutions, the precipitate is obtained in minute orange prisms.

Chloride of potassium (KCl) gives a similar precipitate ($\text{PtCl}_4, 2\text{KCl}$). Platinic chloride having been stated to be a test for potassium and ammonium salts, the reader is prepared to find that potassium and ammonium salts are tests for pla-

tinic salts. The double sodium compound ($\text{PtCl}_4 \cdot 2\text{NaCl}$) is soluble in water.

Collect the precipitate, dry, and heat in a small crucible; it is decomposed, and metal, in the finely divided state of *spongy platinum*, remains.



Heat decomposes the salt of potassium into $\text{Pt} + 2\text{KCl} + \text{Cl}_2$, the chlorine escaping and the chloride of potassium remaining with the platinum.

In working up the platinum residues of laboratory operations, the mixture should be dried, burnt, boiled successively with hydrochloric acid, water, nitric acid, water, then dissolved in aqua regia, excess of acid removed by evaporation, chloride of ammonium added, the precipitate washed with water, dried, ignited, and the resulting spongy platinum retained or converted into perchloride for use as a reagent for alkali-metals. It is by this process that the native platinum is treated to free it from the rare metals palladium, rhodium, osmium, ruthenium, and iridium. The spongy platinum is converted into the massive condition by a refinement on the blacksmith's process of *welding* (German *wellen*, to join), or by fusing in a flame of pure oxygen and hydrogen gases—the oxyhydrogen blow-pipe.

Occlusion by spongy platinum.—Spongy platinum has great power of occlusion. A small piece held in a jet of hydrogen causes ignition of the gas, owing to the close approximation of particles of oxygen (from the air) and hydrogen. Dobereiner's lamp is constructed on this principle—the apparatus being essentially a vessel in which hydrogen is generated by the action of diluted sulphuric acid on zinc, and a cage for holding the spongy platinum.

CADMIUM.

Symbol Cd. Atomic weight 112.

In most of its chemical relations cadmium resembles zinc. In nature it occurs chiefly as an occasional constituent of the ores of that metal. In distilling zinc containing cadmium, the latter, being the more volatile, passes over first. In analytical operations cadmium, unlike zinc, comes down among

the metals precipitated by sulphuretted hydrogen; that is, its sulphide is insoluble in dilute hydrochloric acid, while sulphide of zinc is soluble. It is a white malleable metal, nearly as volatile as mercury. Sp. gr. 8.7.

Beyond the occasional employment of the sulphide as a pigment (*jaune brilliant*), and the iodide in photography and medicine, cadmium and its salts are but little used. The atom of cadmium is bivalent (Cd'').

REACTIONS.

Iodide of Cadmium.

Synthetical Reaction.—Digest together in a flask, metallic cadmium, warm water, and iodine, until the colour of the iodine disappears; solution of iodide of cadmium (*Cadmii Iodidum*, B. P.) (CdI_2) remains. Pearly micaceous crystals may be obtained on evaporating the solution.

This is the process alluded to in the British Pharmacopœia. The compound is used in medicine in the form of ointment, *Unguentum Cadmii Iodidi*, B. P. The salt is also employed with other iodides, in iodizing collodion for photographic purposes. It melts when heated, and is soluble in water or spirit, the solution reddening litmus paper.

First Analytical Reaction.—Through solution of a cadmium salt (CdI_2 or CdCl_2) pass sulphuretted hydrogen; a yellow precipitate of sulphide of cadmium (CdS) falls, resembling in appearance arsenious, arsenic, and stannic sulphides. Add sulphhydrate of ammonium; the precipitate, unlike the sulphides just mentioned, does not dissolve.

Sulphides of cadmium and copper may be separated by solution of cyanide of potassium, in which sulphide of copper is soluble and sulphide of cadmium insoluble.

Second Analytical Reaction.—To a cadmium solution add solution of potash; white hydrate of cadmium (Cd_2HO) is precipitated, insoluble in excess of the potash.

Hydrate of zinc (Zn_2HO), precipitated under similar circumstances, is soluble in solution of potash; the filtrate from

the hydrate of cadmium may therefore be tested for any zinc occurring as an impurity by applying the appropriate reagent—sulphhydrate of ammonium.

Before the blowpipe-flame, on charcoal, cadmium salts give a brown deposit of oxide of cadmium (CdO).

BISMUTH.

Symbol Bi. Atomic weight 208 (Schneider); 210 (Dumas).

Sources.—Bismuth occurs in the metallic state in nature. It is freed from adherent quartz, etc., by simply heating, when the metal melts, runs off, and is collected in appropriate vessels. It is also met with in combination with other elements. Bismuth is greyish white, with a distinct pinkish tinge.

Uses.—Beyond the employment of some of its compounds in medicine, bismuth is but little used. Melted bismuth expands considerably on solidifying, and hence is valuable in taking sharp impressions of dies. It is a constituent of some kinds of type-metal and of pewter-solder.

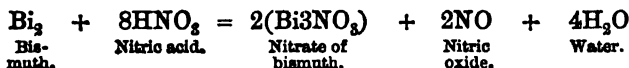
The position of bismuth among the metals is close to that of arsenicum and antimony. Its atom is rarely quinquivalent (Bi^v), but in most compounds trivalent (Biⁱⁱⁱ).

REACTIONS HAVING (a) SYNTHETICAL AND (b) ANALYTICAL INTEREST.

(a) Reactions having Synthetical Interest.

Nitrate of Bismuth.

First Synthetical Reaction.—To a few drops of nitric acid and an equal quantity of water in a test-tube add a little powdered bismuth, heating the mixture if necessary; nitric oxide (NO) escapes and solution of nitrate of bismuth (Bi3NO3) results.



The solution evaporated gives crystals (Bi3NO3, 5H2O), any arsenicum which the bismuth might contain remaining in the mother-liquor. Native bismuth (*Bismuthum*, B. F.) com-

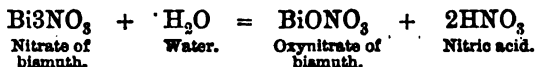
monly contains arsenicum, most of which is removed by roasting or by fusing two or three times with a tenth of its weight of nitre (*Bismuthum Purificatum*, B. P.), or, finally, by converting the metal into oxynitrate, as described in the next reaction, and reducing this with charcoal at a high temperature.

To make *nitrate* of bismuth and other salts on a larger scale, 2 ounces of the metal, in small fragments, are gradually added to a mixture of 4 fluid ounces of nitric acid and 3 of water, and, when effervescence (due to escape of nitric oxide) has ceased, the mixture is heated for ten minutes, poured off from any insoluble matter, evaporated to 2 fluid ounces to remove excess of acid, and then either set aside for *crystals* to form, or poured into half a gallon of water to form the *oxynitrate* of bismuth, or into a solution of 6 ounces of carbonate of ammonium in a quart of water to form the *oxycarbonate* as described in the following reactions.

The precipitates should be washed with cold water and dried at a temperature not exceeding 150° F. Exposed in the moist state to 212° for any length of time, they undergo slight decomposition.

Subnitrate or Oxynitrate of Bismuth.

Second Synthetical Reaction.—Pour some of the above solution of nitrate into a considerable quantity of water; decomposition occurs and oxynitrate of bismuth (BiONO_3) in a hydrous state ($\text{BiONO}_3 \cdot \text{H}_2\text{O}$) (*Bismuthi Subnitratis*, B. P.) is precipitated:—



Filter, and test the filtrate for bismuth by adding excess of carbonate of sodium; a precipitate shows that some bismuth remains in solution. The following equation, therefore, probably more nearly represents the decomposition:—



Decomposition of nitrate of bismuth by water is the process of the British Pharmacopœia for the preparation of oxynitrate

or "subnitrate" of bismuth for use in medicine. For this purpose the original metal must contain no arsenicum. In manufacturing the compound, therefore, before pouring the solution of nitrate into water, the liquid should be tested for arsenicum by one of the hydrogen tests; if that element be present, the solution must be evaporated and only the deposited crystals be used in the preparation of the oxynitrate. For on pouring an arsenical solution of nitrate of bismuth into water, the arsenicum is not wholly removed in the supernatant liquid, unless the oxynitrate be redissolved and reprecipitated several times, according to the amount of arsenicum present.

Subnitrate of bismuth is gradually decomposed by solution of alkaline carbonates; also by the bicarbonates with production of carbonic acid gas; oxycarbonate of bismuth and nitrate of the alkali-metal being formed. It is sometimes administered in the form of a lozenge (*Trochisci Bismuthi*, B. P.). It is also used as a cosmetic under the name of *Pearl-white* (*Blanc de Perle*).

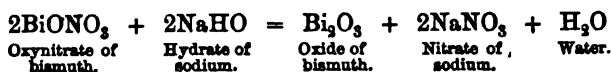
Oxysalts of Bismuth.—It will be noticed that the formula for subnitrate of bismuth (BiNO_3) does not accord with that of other nitrates, the characteristic elements of which are NO_3 . Analogy would seem to indicate, however, that the fourth atom of oxygen has different functions to the three in the NO_3 ; for on pouring solution of chloride of bismuth (BiCl_3) into water, oxychloride is produced (BiOCl) (a white powder used as a cosmetic, also in enamels and in some varieties of sealing-wax). The bromide (BiBr_3) and iodide (BiI_3) similarly treated yield oxybromide (BiOBr) and oxyiodide (BiOI). The subnitrate (BiNO_3) is, therefore, probably an analogous compound, an oxynitrate (BiONO_3). The sulphate (Bi_2SO_4) also decomposes when placed in water, giving what may be termed an oxysulphate, the formula of which is $\text{Bi}_2\text{O}_5\text{SO}_4$.

It is difficult to prove whether or not the water in the "subnitrate" or hydrous oxynitrate of bismuth ($\text{BiONO}_3 \cdot \text{H}_2\text{O}$) is an integral part of the salt. If it is, the compound is simply the hydrato-nitrate ($\text{BiNO}_3 \cdot 2\text{HO}$) of bismuth.

Oxide of Bismuth.

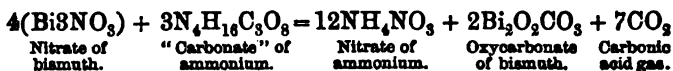
Third Synthetical Reaction.—Boil subnitrate of bismuth with solution of soda for a few minutes; it is converted

into yellowish oxide of bismuth (Bi_2O_3) (*Bismuthi Oxidum*, B. P.).



Subcarbonate or Oxycarbonate of Bismuth.

Fourth Synthetical Reaction.—To solution of nitrate of bismuth add carbonate of ammonium; a white precipitate of hydrous oxycarbonate ($2\text{Bi}_2\text{O}_3\text{CO}_3, \text{H}_2\text{O}$) (*Bismuthi Carbonas*, B. P.) falls.



This compound may be regarded as similar in constitution to the oxysalts just described. In Bi_2CO_3 one scarcely recognises the characteristic elements of carbonates; but considering the preparation to be an oxycarbonate ($\text{Bi}_2\text{O}_3\text{CO}_3$) its relations to carbonates and oxides are evident. These subsalts may all be viewed as normal bismuth salts in which an atom of oxygen displaces an equivalent proportion of other acidulous atoms or radicals:—

Chloride	Bi_3Cl	Oxychloride . . .	BiOCl
Bromide	Bi_3Br	Oxybromide . . .	BiOBr
Iodide	Bi_3I	Oxyiodide . . .	BiOI
Nitrate	Bi_3NO_3	Oxynitrate . . .	BiONO_3
Sulphate	Bi_3SO_4	Oxysulphate . . .	$\text{Bi}_2\text{O}_3\text{SO}_4$
Carbonate (unknown)	Bi_3CO_3	Oxycarbonate . . .	$\text{Bi}_2\text{O}_3\text{CO}_3$

They may be viewed, in short, as salts in process of conversion to oxide; continue the substitution a little further, and each yields oxide of bismuth (Bi_2O_3). They have also been considered to be salts of a hypothetical univalent radical bismuthyl (BiO).

Citrate of Bismuth.

Fifth Synthetical Reaction.—To solution of nitrate of bismuth add citric acid and then solution of ammonia until the precipitate at first formed is redissolved, and the liquid after

shaking has a slight ammoniacal odour. The product contains citrate of bismuth ($\text{BiC}_6\text{H}_5\text{O}_7$), dissolved in solution of citrate and nitrate of ammonium. Made with definite quantities of ingredients and an amount of bismuth salt equivalent to three grains of oxide (Bi_2O_3) in a fluid drachm, the solution forms the *Liquor Bismuthi et Ammoniac Citratis*, B. P.).

If the ordinary oxynitrate of bismuth be heated for a few moments with a concentrated solution of an equivalent of citric acid, the normal citrate is generated as a heavy crystalline powder, and the nitric acid entirely freed and easily washed away by decantation. Ten parts of the oxynitrate, 7 of citric acid crystals, and 30 to 40 of water are heated together for a few minutes, until a drop of the mixture forms a clear solution with ammonia water; the crystalline mass is next diluted with eight to ten times its volume of water, and set aside for a short time to let the citrate deposit; the clear liquid is afterwards decanted. The crystalline sediment is then washed three or four times in a similar manner, drained, and either dried on a water-bath or by mere exposure. The yield is about 13 2-3 parts, showing that the salt is anhydrous, and that its formula is $\text{Bi}''\text{C}_6\text{H}_5\text{O}_7$ (Rother).

(b) *Reactions having Analytical Interest (Tests).*

First Analytical Reaction.—Through solution of a bismuth salt (a slightly acid solution of nitrate, for example) pass sulphuretted hydrogen; a black precipitate of sulphide of bismuth (Bi_2S_3) falls. Add ammonia (to neutralize acid) and then sulphhydrate of ammonium; the precipitate, unlike As_2S_3 and Sb_2S_3 , is insoluble.

Second Analytical Reaction.—Concentrate almost any acid solution of a bismuth salt and pour into water; a white salt is precipitated.

This reaction is characteristic of bismuth salts: it has already been amply explained. The precipitate is distinguished from one formed by antimony under similar circumstances, by being insoluble in solution of tartaric acid.

Third Analytical Reaction.—To a solution of bismuth salt

add an alkali; hydrate of bismuth (Bi_3HO) is precipitated insoluble in excess.

Fourth Analytical Reaction.—A small quantity of the following reagent, including both supernatant liquid and precipitated yellow scales, is transferred to a test-tube, and gradually heated till solution takes place. Any liquid containing, or supposed to contain, bismuth is then added, and the whole allowed to cool. The separated scales will show a distinct change in colour to dark orange or crimson according to the quantity of bismuth present.

The test liquid may be prepared by adding to a boiling solution of acetate of lead, containing about $\frac{1}{2}$ gr. to the ounce, solution of iodide of potassium in considerable excess. The solution of the iodide of lead precipitated is assisted by a little acetic acid. On cooling, iodide of lead is deposited in the characteristic yellow crystalline plates or scales.

The reader is again advised to trace out the exact nature of each of the foregoing reactions, chiefly by aid of equations or diagrams.

QUESTIONS AND EXERCISES.

341. Enumerate the fifteen metals, salts of which are frequently employed in pharmacy.
342. Mention the twelve rarer metals interesting to pharmacists.
343. Name the sources and official compounds of lithium.
344. Explain the formation of Citrate of Lithium.
345. What is the strength of *Liquor Lithias Effervescens*?
346. On what chemical hypothesis are lithium compounds administered to gouty patients?
347. Describe the relation of lithium to other metals.
348. What is the chief test for lithium?
349. Write a paragraph on strontium, its natural compounds, chemical relations, technical applications, and tests.
350. What are the formula and properties of oxalate of cerium?
351. Name the commonest ore of manganese; and give an equation descriptive of its reaction with hydrochloric acid.

352. Explain the formation of permanganate of potassium, employing diagrams or equations.

353. In what manner do the manganates of potassium act as disinfectants?

354. What are the chief tests for manganese?

355. What are the chief uses of the compounds of cobalt?

356. How are salts of cobalt analytically distinguished from those of nickel?

357. Mention an application of nickel in the arts.

358. What is the general colour of nickel salts?

359. State the method of preparation of red chromate of potassium.

360. Give the formulæ of red and yellow chromates of potassium.

361. How is red chromate of potassium obtained?

362. Describe the action of sulphuretted hydrogen on acidified solutions of chromates.

363. What is the formula of chrome alum?

364. Mention the chief tests for the chromic radical, and for chromium.

365. How would you detect iron, chromium, and aluminium in a solution?

366. Define the terms tinstone, stream-tin, block tin, grain tin, tin-plate.

367. Describe the position occupied by tin in relation to other metals.

368. What is the difference between stannic acid and meta-stannic acid?

369. State the applications of tin in the arts.

370. Mention the chief tests for stannous and stannic salts.

371. Name the best antidote in cases of poisoning by tin solutions.

372. How is gold dust separated from the earthy matter with which it is naturally associated?

373. How much pure gold is contained in English coin and in jewellers' gold?

374. State the average thickness of gold leaf.

375. What is the weight of a sovereign?

376. Explain the term "fineness" as applied to gold.

377. What effect is produced on gold by hydrochloric, nitric, and nitrohydrochloric acids respectively?

378. By what reagent is metallic gold precipitated from solutions of its salts?
379. How is Purple of Cassius prepared?
380. Whence is platinum obtained?
381. Why are platinum utensils peculiarly adapted for use in chemical laboratories?
382. How is perchloride of platinum prepared?
383. Name the chief tests for platinum.
384. What is "platinum black"?
385. Describe an experiment demonstrative of the large amount of attraction for gases possessed by metallic platinum.
386. How is "spongy platinum" produced?
387. By what process may the metal be recovered from platinum residues?
388. What is *occlusion* in chemistry?
389. In what condition does cadmium occur in nature?
390. By what process may Iodide of Cadmium be prepared? and in what form is it used in medicine?
391. Mention the chief test for cadmium.
392. Distinguish sulphide of cadmium from other sulphides of similar colour.
393. How is cadmium separated from zinc?
394. How does bismuth occur in nature?
395. What is the quantivalence of bismuth?
396. Write down equations descriptive of the action of nitric acid on bismuth, and water on nitrate of bismuth.
397. How may pure salts be prepared from bismuth containing arsenicum?
398. Give a diagram of the official process for the so-called Carbonate of Bismuth.
399. Write formulæ showing the accordance of the official Subnitrate and Carbonate with the other salts of Bismuth, and with ordinary Nitrates and Carbonates.
400. How is *Liquor Bismuthi et Ammoniac Citratis* prepared?
401. What are the tests for Bismuth?
-

Practical Analysis.

Bismuth is the last of the metals whose synthetical or analytical relations are of general interest. The position of the rarer among the common metals, and the influence which either has on the other during the manipulations of analysis, will now be considered. These objects will be best accomplished, and a more intimate acquaintance with all the metals be obtained, by analysing, or studying the methods of analysing, solutions containing one or more metallic salts.

Of the following Tables, the first includes directions for the analysis of an aqueous or only slightly acid solution containing but one salt of any of the metals hitherto considered. Here the colour of the precipitate or precipitates afforded by a metal under given circumstances must largely be relied on in attempting the detection of the various elements.

The long, folded Table is intended as a chart for the analysis of solutions containing salts of more than one of the common and rarer metals. It is simply a compilation from the foregoing reactions—an extension of the scheme for the analysis of salts of the ordinary metals. Hence it often may be altered or varied in arrangement to suit the requirements of the analyst.

That on p. 297 is a mere outline of the other two Tables. It gives the position of the metals in relation to each other, and will much aid the memory in recollecting that relation.

The analysis of solutions containing only one metal will, as already stated, serve to impress the memory with the characteristic tests for the various metals and other radicals, and familiarize the mind with chemical principles. Medical students seldom have time to go further than this. More thorough analytical and general chemical knowledge is only acquired by working on such mixtures of bodies as are met with in actual practice, beginning with solutions which may contain any or all of the members of a group (*see* previous pages), then examining solutions containing more than one group, and finally analysing liquids in which are dissolved several salts of any of the common or rarer metals.

The Author cannot too strongly recommend students thoroughly to master the art of analysis, not only on account of its direct value, but because its practice enables the learner rapidly and soundly to acquire a good knowledge of chemistry, and greatly to improve his general mental faculties.

TABLE OF SHORT DIRECTIONS FOR THE ANALYSIS OF AN AQUEOUS OR ONLY SLIGHTLY ACID SOLUTION OF ORDINARY SALTS OF ONE OF THE ELEMENTS HITHERTO CONSIDERED—THE COMMON AND RARE METALS.

Add hydrochloric acid.

Precipitate Hg (ous) Pb Ag. Collect, wash, and add AmHO. Hg ppt., blackened. Pb ppt., still white. Ag ppt., dissolved. Sb and Bi may also be precipitated by HCl, but are dissolved on adding more HCl.	If HCl gave no precipitate the metal is still in the liquid; pass H ₂ S through the solution.		
	If H ₂ S gave no precipitate add AmCl, AmHO, and AmHS.		
Precipitate Cd Cu Hg(ic) Pb Bi As Sb Sn Au Pt. Collect, wash, add AmHS. Insoluble. Cd, yellow Cu Hg(ic) Pb Bi	Precipitate Zn Mn Co Ni Al Fe Or. Zn } white. Al } Cr, green. Mn, skin-tint. Ni } Co } black. Fe }	If AmHS gave no precipitate add Am ₂ CO ₃ . Precipitate Ba Sr Ca. Collect, wash, dissolve in HCl, H ₂ O ₂ , add K ₂ CrO ₄ . Ppt. Sol. Ba. Sr Ca. Add dil. H ₂ SO ₄ . Ppt. Sol. Sr. — Ca.	If Am ₂ CO ₃ gave no precipitate add Am ₂ HAsO ₄ . Ppt. Mg. If no precipitate, test original solution in flame on loop of Pt. wire. L, crimson Na, yellow K, violet. If neither, test orig. sol. for Am.
Apply special tests for each to the original solution. For these, see the previous pages.	Test specially for each in original solution. See previous pages.		

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OUTLINE OF THE ANNEXED ANALYTICAL TABLES.

HCl	H ₂ S	AmHS	Am ₂ CO ₃	Am ₂ HAsO ₄	
Hg (as mercurous salt)	Cd	Zn	Ba	Mg	K
	Cu	Mn*	Sr		Na
Pb (partially)	Hg (as mercuric salt)	Co	Ca		Am
	Pb (entirely)	Ni			
Ag	Bi	Al		L	
	As (as arsenious or arsenic salt)	Fe			
	Sb	Cr			
	Sn (as stannous or stannic salt)				
	Au				
	Pt				

The laboratory student should practise the examination of aqueous solutions of salts of the above metals until he is able to analyse with facility and accuracy.

* See p. 265.

GENERAL AND SPECIAL MEMORANDA RELATING TO THE
PRECEDING ANALYTICAL TABLES.

General Memoranda.

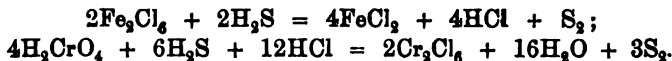
These charts are constructed for the analysis of salts more or less soluble in water.—The student has still to learn how substances insoluble in water are to be brought into a state of solution: but, once dissolved, their analysis is effected by the same scheme as that just given. The Tables, especially the longer, folded one, may therefore be regarded as fairly representing the method by which metallic constituents of chemical substances are separated from each other and recognised.—The methods of isolation of the complementary constituent of the salt (the reactions of non-metals and acidulous radicals) will form the next object of practical study.

The general memoranda given in connection with the first Table (p. 253) are equally applicable to the extended second Table, and should again be carefully read through.

Special Memoranda.

The hydrochloric-acid precipitate may at first include some antimony and bismuth as oxychlorides, readily dissolved, however, by excess of acid.—If either of these elements be present the washings of the precipitate will probably be milky; in that case add a few drops of hydrochloric acid, which will clear the liquid and make way for the application of the test for lead.

The sulphuretted-hydrogen precipitate may be white, in which case it is nothing but sulphur; for, as already indicated, ferric salts are reduced to ferrous, and chromates to the lower salts of chromium by sulphuretted hydrogen, sulphur being deposited:—



But the precipitate may also be coloured, or even white when only lead or mercury are present, through an insufficiency of sulphuretted hydrogen having produced a peculiar oxysulphide or hydrato-sulphide, etc. The gas should be

passed through the liquid until, even after well shaking, it smells strongly of sulphuretted hydrogen.

The portion of the sulphuretted-hydrogen precipitate dissolved by sulphhydrate of ammonium may include a trace of copper, sulphide of copper being not altogether insoluble in sulphhydrate of ammonium.——On adding hydrochloric acid to the sulphhydrate-of-ammonium solution, a white precipitate of sulphur only may be precipitated, the sulphhydrate of ammonium nearly always containing free sulphur.——Strong hydrochloric acid does not readily dissolve small quantities of sulphide of antimony out of much sulphide of arsenicum; and, on the other hand, the strong hydrochloric acid takes into solution a small quantity of sulphide of arsenicum if much sulphide of antimony is present. The precipitates or the original solutions should therefore be examined by the other (hydrogen) tests for these elements if doubt exists concerning the presence or absence of either. Tin remains in the hydrogen-bottle in the metallic state, deposited as a black powder on the zinc used in the experiment. The contents of the bottle are turned out into a dish, ebullition continued until evolution of hydrogen ceases, and the zinc is taken up by the excess of sulphuric acid employed; any tin is then filtered out, washed, dissolved in a few drops of hydrochloric acid, and the liquid tested for tin by the usual reagents.——Tin may be detected in the mixed sulphides of tin, arsenicum, and antimony by the blow-pipe reaction (p. 278).

The portion of the sulphuretted-hydrogen precipitate not dissolved by the sulphhydrate of ammonium may leave a yellow semi-fused globule of sulphur on boiling with nitric acid. This globule may be black, not only from presence of mercuric sulphide, but also from enclosed particles of other sulphides protected by the sulphur from the action of the acid. It may also contain sulphate of lead, produced by the action of nitric acid on sulphide of lead. In cases of doubt the mass must be removed from the liquid, boiled with nitric acid till dissolved, the solution evaporated to remove excess of acid, and the residue examined; but usually it may be disregarded.——Before testing for bismuth, any considerable excess of acid should be removed by evaporation, and the residual liquid should be freely diluted. If no precipitate (oxynitrate of bismuth) appear, chloride-of-ammonium solution may be added, oxychloride of bismuth more readily forming than even oxynitrate.——Or any nitric acid or sulphuric acid having been neutralized.

by ammonia, hydrochloric acid is added and then iodide of potassium; a rich orange colour results if bismuth be present. In testing for lead by sulphuric acid the liquid should be diluted and set aside for some time.

Mercury may also be isolated by digesting the sulphuretted-hydrogen precipitate in sulphhydrate of sodium instead of sulphhydrate of ammonium. The sulphides of arsenicum, antimony, tin, and mercury are thus dissolved out. The mixture is then filtered, excess of hydrochloric acid added to the filtrate, and the precipitated sulphides collected on a filter, washed, and digested in sulphhydrate of ammonium; sulphide of mercury remains insoluble, while the sulphides of arsenicum, antimony, and tin are dissolved. By this method copper also appears in its right place only, sulphide of copper being quite insoluble in sulphhydrate of sodium. The other metals are then separated in the usual way.

The sulphhydrate-of-ammonium precipitate may, if the original solution was acid, contain Phosphates, Oxalates, Silicates, and Borates of Barium, Calcium, and Magnesium. These will subsequently come out with the iron, and, being white, give the iron precipitate a light-coloured appearance; their examination must be conducted separately, by a method described subsequently in connection with the treatment of substances insoluble in water.—The precipitate containing aluminium, iron, and chromium hydrates often contains some manganese. This manganese may be detected by washing the hydrates to remove all trace of chlorides, boiling with nitric acid, adding either puce-coloured oxide of lead or red lead, and setting the vessel aside; if manganese be present a red or purple liquid is produced.—Sulphide of nickel is not easily removed by filtration (*vide* p. 269) until most of the excess of sulphhydrate of ammonium has been dissipated by prolonged ebullition.

The carbonate-of-ammonium precipitate may not contain the whole of the barium, strontium, and calcium in the mixture, unless free ammonia be present; for the carbonates of those metals are soluble in water charged with carbonic acid. If, therefore, the liquid is not distinctly ammoniacal, solution of ammonia should be added.—Neither carbonate nor hydrate of ammonium wholly precipitates magnesian salts; and as a partial precipitation is undesirable, a solvent, in the form of an alkaline salt (chloride of ammonium), if not already in the liquid, should be added.—In the chart opposite p. 296 strontium is ordered to be separated from calcium by adding

to the acetic solution diluted sulphuric acid. The latter, unless extremely dilute, may precipitate calcium. Any such loss of calcium is in itself of little consequence, because enough sulphate of calcium remains in the filtrate to afford a calcium reaction when ammonia and oxalate of ammonium are subsequently added. But the calcium precipitated by the sulphuric acid may be wrongly set down as strontium. Therefore test a little of the acetic solution for strontium by an aqueous solution of sulphate of calcium, when, if no precipitate falls after setting aside for several minutes, strontium may be regarded as absent. If a precipitate occurs strontium is present; the rest of the acetic solution is then tested for calcium as directed in the chart, the final testing by oxalate of ammonium being, of course, preceded by the addition of ammonia.

Lithium.—The search for lithium may usually be omitted. Should a precipitate, supposed to be due to lithium, be obtained, it must be tested in a flame (= scarlet tint), as a little magnesium not unfrequently shows itself under similar circumstances.

Spectral Analysis.—If present only in minute proportions, the lithium may also remain with the alkalis; it can then only be detected by physical analysis (by a prism) of the light emitted from a tinged flame—by, in short, an instrument termed a spectroscope. Such a method of examination is called spectral analysis, a subject of much interest and of no great difficulty, but scarcely within the range of Pharmaceutical Chemistry; it will be briefly described in connection with the methods of analysing solid substances.

QUESTIONS AND EXERCISES.

402. Describe a general method of analysis by which the metal of a single salt in a solution could be quickly detected.

403. Give illustrations of black, white, light pink, yellow, and orange sulphides.

404. Mention the group-tests generally employed in analysis.

405. Under what circumstances may a hydrochloric precipitate contain antimony or bismuth?

406. If a sulphuretted-hydrogen precipitate is white, what substances are indicated?

407. Give processes for the qualitative analysis of liquids containing the following substances:—

- a. Arsenicum and Cadmium.
- b. Bismuth and Antimony.
- c. Ferrous and Ferric salts.
- d. Aluminium, Iron, and Chromium.
- e. Arsenicum, Antimony, and Tin.
- f. Lead and Strontium.
- g. Iron, Sodium, and Arsenicum.
- h. Mercury, Manganese, and Magnesium.
- i. Zinc, Manganese, Nickel, and Cobalt.
- j. Barium, Strontium, and Calcium.

THE ACIDULOUS RADICALS.

Introduction.—The twenty-seven radicals which have up to this point mainly occupied attention are (admitting ammonium, NH_4) metals; and they have been almost exclusively studied not in the free state, but in the condition in which they exist in salts. Moreover these metals have been treated as if they formed the more important constituent, the stronger half, the foundation or *base* of salts. Attention has been continuously directed to the metallic or *basyulous* side of salts. And, indeed, there is still one more basyulous radical which must be mentioned, though it is usually supposed to play only a subordinate part in medicine—Hydrogen. Unlike the salts of most metals, those of hydrogen (the so-called *acids*) are never, in medicine or the arts generally, professedly used for the sake of their hydrogen, but always for the other half of the salt, the *acidulous* side. And it is not for their basyulous radical that these hydrogen salts are now commended to notice,* but in order to study, under the most favourable cir-

* It must not be forgotten that the commonest salt of any radical whatsoever is a salt of hydrogen, the oxide of hydrogen (H_2O), or hydrate of hydrogen (HHO), *water*. In the reactions already performed, the value of this compound has been constantly recognised, both for its hydrogen and for its oxygen, but most of all as the vehicle or medium by which nearly all other atoms are enabled to come into that contact with each other without which their existence would be almost useless; for

cumstances, those acidulous groupings which have continually presented themselves in operations on salts, but which were for the time of secondary importance. These *acidulous radicals* may now be treated as the primary object of attention; and there is no better way of doing so than by operating on their compounds with hydrogen, the relatively inferior medicinal importance of which element, as compared with potassium, iron, and other basylous radicals, will serve to give the desired prominence to the *acidulous radicals* in question.

Common Acids.—These salts of hydrogen (hydrogen easily displaceable, wholly or, in certain cases, in part, by ordinary metals) are the ordinary, sharp, sour bodies termed *acids* (from the Latin root *acies*, an edge). The following Table includes the formulæ and usual names of the most important: others will be noticed subsequently. A few of those mentioned are unstable or somewhat rare; in such cases a common metallic salt containing the acidulous radical may be used for reactions.

HCl . . .	hydrochloric acid.
HBr . . .	hydrobromic acid.
HI . . .	hydriodic acid.
HCN (HCy)	hydrocyanic acid.
HNO ₃ . . .	nitric acid.
HClO ₃ . . .	chloric acid.
HC ₂ H ₃ O ₂ . .	acetic acid.*
H ₂ S . . .	hydrosulphuric acid.†
H ₂ SO ₃ . . .	sulphurous acid.
H ₂ SO ₄ . . .	sulphuric acid.

some atoms are like some animals, out of water they are as inactive as fishes. It is true that both fishes and salts have usually to be removed from water to be utilized by man; but before they can be assimilated, either as food or as medicine, they must again seek the agency of water—in becoming dissolved.

* The hydrogen on the acidulous side must not be confounded with the basylous hydrogen in all these hydrogen salts or acids; the two perform entirely different functions. Hydrogen in the acidulous portion is like the hydrogen in the basylous radical ammonium; it has combined with other atoms, to form a group which plays more or less the part of an elementary radical, and to which a single symbol is not unfrequently applied (Am, Cy, A, O, T, C, etc.). Cobalt, chromium, iron, platinum, etc., resemble hydrogen in this respect in often uniting with other atoms to form definite acidulous radicals in which the usual basylous character of the metals has for the time disappeared. In *hydrides* (p. 133) hydrogen itself is an acidulous radical.

† Synonyms:—sulphydric acid and sulphuretted hydrogen.

H_2CO_3 ?	. carbonic acid.
$\text{H}_2\text{C}_2\text{O}_4$. .	oxalic acid.
$\text{H}_3\text{C}_4\text{H}_4\text{O}_6$.	tartaric acid.
$\text{H}_3\text{C}_6\text{H}_5\text{O}_7$.	citric acid.
H_3PO_4 . .	phosphoric acid.
H_3BO_3 . .	boracic acid.

The old names are here retained for these acids; but in studying their chemistry and chemical relations to other salts they are usefully spoken of by such more purely chemical names as (for hydrochloric acid) chloride of hydrogen, (for nitric acid) nitrate of hydrogen, and so on—sulphate of hydrogen, tartrate of hydrogen, phosphate of hydrogen.

A prominent point of difference will at once be noticed between the basylous radicals met with up to the present time and the acidulous groupings included in the above tabular list. The former are nearly all elements, ammonium only being a compound; the latter are mostly compounds, chlorine, bromine, iodine, and sulphur being the only elements. This difference will not, however, be so apparent when the chemistry of alcohols, ethers, and such bodies has been mastered; for they are all salts of *compound* basylous radicals.

Rarer Acids.—The above acids contain the only acidulous groupings that commonly present themselves in analysis or in pharmaceutical operations. There are, however, several other acids (such as hypochlorous, nitrous, hypophosphorous, valerianic, benzoic, gallic, tannic, uric, hyposulphurous, hydroferrocyanic, hydroferridcyanic, lactic, etc.) with which it is desirable to be more or less familiar; reactions concerning these will therefore be described. Arsenious, arsenic, stannic, manganic, and chromic acids have already been treated of in connection with the metals they contain; in practical analysis these acids always become sufficiently altered for their metals to come out among the basylous radicals.

Quantivalence.—A glance at the foregoing table is sufficient to show the quantivalence of the acidulous radicals. The first seven are clearly univalent; then follow six bivalent, leaving three trivalent.

These all combine with equivalent amounts of basylous radicals to form various salts; hence they may be termed monobasylous, dibasylous, and tribasylous radicals. The acids themselves were formerly spoken of as monobasic, dibasic, and tribasic respectively, or monobasic and polybasic,

in reference to the amount of *base* (hydrates or oxides) they could decompose; but the terms are no longer definite, and hence but little used in mineral chemistry.

Antidotes.—The antidotes in cases of poisoning by the strong acids will obviously be non-corrosive alkaline substances, as soap and water, magnesia, and common washing "soda" or other carbonates. Vinegar, lemon-juice, and weak or non-corrosive acids would be the appropriate antidotes to caustic alkalies.

Analysis.—The practical study of the acidulous side of salts will occupy far less time than the basylous. Salts will then be briefly examined as a whole.

One word of caution.—It is only for convenience in the division of chemistry for systematic study that salts may be considered to contain basylous and acidulous radicals, or separate sides, so to speak; for we possess no absolute knowledge of the internal arrangement of the atoms (admitting that there are such things) in the molecule of the salt. We only know that certain groups of atoms may be transferred from compound to compound in mass (that is, without apparent decomposition); hence the assumption that these groups are radicals. A salt is probably, however, a whole, having no such sides as those mentioned.

QUESTIONS AND EXERCISES.

408. Mention the basylous radical of acids.

409. Give illustrations of univalent, bivalent, and trivalent acidulous radicals, or monobasylous, dibasylous, and tribasylous radicals.

410. What is the difference between an elementary and a compound acidulous radical?

411. Name the grounds on which salts may be assumed to contain basylous and acidulous radicals.

HYDROCHLORIC ACID AND OTHER CHLORIDES.

Formula of Hydrochloric Acid HCl . Molecular weight* 36.5.

The acidulous radical of hydrochloric acid and of other

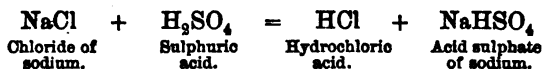
* The weight of a molecule is the sum of the weights of its atoms.

chlorides is the element chlorine (Cl). It occurs in nature chiefly as chloride of sodium (NaCl), either solid, under the name of *rock-salt*, mines of which are not unfrequently met with, or in solution in the water of all seas. Common table-salt is more or less pure chloride of sodium in minute crystals. Chlorine, like hydrogen, is univalent (Cl'); its atomic weight is 35.5. Its molecule is symbolized thus, Cl₂, chloride of chlorine.

REACTIONS.

Hydrochloric Acid.

First Synthetical Reaction.—To a few fragments of chloride of sodium in a test-tube or small flask add about an equal weight of sulphuric acid; colourless and invisible gaseous hydrochloric acid is evolved, a sulphate of sodium remaining. Adapt to the mouth of the vessel, by a perforated cork, a piece of glass tubing bent to a right angle, heat the mixture, and convey the gas into a small bottle containing a little water; solution of hydrochloric acid results.

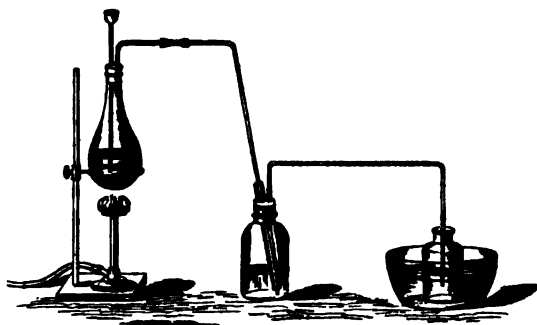


Hydrochloric acid.—The product of this operation is the nearly colourless and very sour liquid commonly termed hydrochloric acid. When of certain given strengths (estimated by volumetric analysis) it forms *Acidum Hydrochloricum*, B. P., and *Acidum Hydrochloricum Dilutum*, B. P. The former has a specific gravity of 1.16 (1.1578), and contains 31.8 per cent. of real acid; the latter, sp. gr. 1.052, with 10.58 per cent. of the real acid, is made by diluting 8 fluid parts of the strong acid with water until the mixture measures 26½ fluid parts. The above process is that of the Pharmacopœia—larger vessels being employed, and the gas being freed from any trace of sulphuric acid by washing. Other chlorides yield hydrochloric acid when heated with sulphuric acid; but chloride of sodium is always used because cheap and common.

Common yellow hydrochloric acid is a by-product in the manufacture of carbonate of sodium from common salt, a process in which the chloride of sodium is first converted into sulphate, hydrochloric acid being liberated. This impure acid is liable to contain iron, arsenic, fixed salts,

sulphuric acid, sulphurous acid, nitrous compounds, and chlorine.

Fig. 87.



PREPARATION OF HYDROCHLORIC ACID.

The official process for the preparation of hydrochloric acid is as follows: it may be carried out by the student with about one-twelfth of the quantities mentioned:—

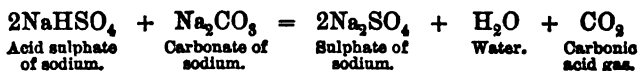
“Take of chloride of sodium, dried, 48 ounces, sulphuric acid 44 fluid ounces, water 36 fluid ounces, distilled water 50 fluid ounces. Pour the sulphuric acid slowly into thirty-two ounces of the water, and, when the mixture has cooled, add it to the chloride of sodium previously introduced into a flask having the capacity of at least one gallon. Connect the flask by corks and a bent glass tube with a three-necked wash-bottle, furnished with a safety-tube, and containing the remaining four ounces of the water (or let the flask-tube pass loosely through a wider tube fixed in the cork of the wash-bottle, as shown in fig. 87); then, applying heat to the flask, conduct the disengaged gas through the wash-bottle into a second bottle containing the distilled water, by means of a bent tube dipping about half an inch below the surface, and let the process be continued until the product measures sixty-six ounces, or the liquid has acquired a specific gravity of 1.16. The bottle containing the distilled water must be kept cool during the whole operation.”

The modification of wash-bottle shown in the figure allows of the easy insertion or removal of a delivery-tube. The wider tube there shown, or an ordinary tube-funnel, also acts

as a *safety-tube* by admitting air the moment there is any tendency in the water in the receiver to be forced back on account of a too rapid absorption of the gas. The time of the student will be saved if hydrochloric acid already in stock be placed in the wash-bottle instead of water.

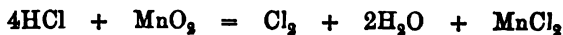
Invisible gaseous hydrochloric acid forms visible greyish-white fumes on coming into contact with air. This is due to combination with the moisture of the air. The intense greediness of hydrochloric gas and water for each other is strikingly demonstrated on opening a test-tube full of the gas under water; the latter rushes into and instantly fills the tube. If the water is tinged with blue litmus, the acid character of the gas is prettily shown at the same time. The test-tube, which should be perfectly dry, may be filled from the delivery-tube direct; for the gas is somewhat heavier than, and therefore readily displaces, air. The mouth may be closed by the thumb of the operator. At low temperatures hydrochloric acid and water form a crystalline compound $\text{HCl} \cdot 2\text{H}_2\text{O}$.

Note.—The process, as described in the British Pharmacopœia, includes the use of as much sulphuric acid as is theoretically necessary for the production of acid sulphate of sodium (NaHSO_4) which remains in the generating vessel. A hot solution of this residue, carefully neutralized by carbonate of sodium, filtered and set aside, yields normal sulphate (*Sodæ Sulphas*, B. P.) in the form of transparent oblique efflorescent prisms ($\text{Na}_2\text{SO}_4 \cdot 10\text{H}_2\text{O}$).



Chlorine.

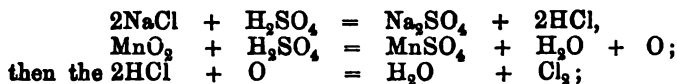
Second Synthetical Reaction.—To some drops of hydrochloric acid (that is, the common aqueous solution of the gas) add a few grains of black oxide of manganese, and warm the mixture; *chlorine*, the acidulous radical of all chlorides, is evolved, and may be recognised by its peculiar odour, or irritating effect on the nose and air-passages.



Chlorine-water.—This is the process of the British Pharma-

copœia for the production of chlorine-water (*Liquor Chlori*, B. P.), the gas being first washed and then passed into water. One ounce of oxide to six fluid ounces of acid diluted with two of water, and the gas passed through a wash-bottle containing about two ounces of water, yield enough chlorine to produce about a pint and a half of chlorine-water. (On this small scale, less than half the acid is utilized through incomplete decomposition of the materials and, especially, through incomplete absorption of the chlorine gas.) Chlorine slowly decomposes water with production of hydrochloric acid and oxygen gas; it is best preserved in a green-glass well-stoppered bottle in a cool and dark place. At common temperatures (60° F.), if fresh and thoroughly saturated, chlorine-water contains more than twice (2·3) its bulk of chlorine, or less than 1 per cent. (about 0·75) by weight. Chlorine passed into cold water yields crystals of hydrous chlorine ($\text{Cl}_5\text{H}_2\text{O}$), and these, when heated under pressure, give an upper layer of chlorine-water and a lower layer of *liquid chlorine*.

Note.—To obtain the chlorine from other chlorides, sulphuric acid, as well as black oxide of manganese, must be added. Hydrochloric acid is first formed. The action described in the foregoing equation then goes on, except that half instead of the whole of the oxygen from the black oxide is available for the removal of the hydrogen from the chlorine of the hydrochloric acid, the other half being taken up by the hydrogen of the sulphuric acid. Thus, supposing common salt to be the chloride used, the following equations may represent the supposed steps of the process:—



or the whole may be included in one equation:



This reaction may have occasioned analytical interest, a very small quantity of combined chlorine being recognised by its means. But the following test is nearly always applicable for the detection of this element, and leaves nothing to be desired in point of delicacy.

Analytical Reactions (Tests).

To a drop of hydrochloric acid, or to a dilute solution of any other chloride, add solution of nitrate of silver; a white curdy precipitate falls. Pour off most of the supernatant liquid, add nitric acid, and boil; the precipitate does not dissolve. Pour off the acid, and add dilute ammonia; the precipitate quickly dissolves. Neutralize the solution by an acid; chloride of silver is once more precipitated.

The formation of this white precipitate, its appearance, insolubility in boiling nitric acid, solubility in ammonia, and reprecipitation by an acid form abundant evidence of the presence of chlorine. Its occurrence as a chloride of a metal is determined by testing for the metal with the appropriate reagents; its occurrence as hydrochloric acid is considered to be indicated by the odour, if strong, and the sour taste, if weak, of the liquid, and the action of the liquid on blue litmus paper, which, like other acids, it reddens. If hydrochloric acid be present in excessive quantity it will, in addition to the above reactions, give rise to strong effervescence on the addition of a carbonate, a chloride being formed. The chlorine in insoluble chlorides, such as calomel, "white precipitate," etc., may be detected by boiling with caustic potash, filtering, acidulating the filtrate by nitric acid, and then adding the nitrate of silver.

Antidotes.—In cases of poisoning by strong hydrochloric acid, solution of carbonate of sodium (common washing-soda) or a mixture of magnesia and water may be administered as an antidote.

QUESTIONS AND EXERCISES.

412. The official Hydrochloric Acid contains 31·8 per cent. by weight of gas, and its specific gravity is 1·16; work out a sum showing what volume of it will be required, theoretically, to mix with black oxide of manganese for the production of one gallon of chlorine-water, one fluid ounce of which contains 2·66 grains of chlorine. Ans. $5\frac{1}{2}$ fl. ozs., nearly (5·4).

413. Why does hydrochloric acid gas give visible fumes on coming into contact with air?

414. How much chloride of sodium will be required to furnish one pound of chlorine?

415. Give the analytical reactions of chlorides.

416. What antidotes may be administered in cases of poisoning by hydrochloric acid?

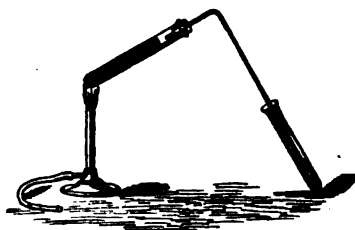
HYDROBROMIC ACID AND OTHER BROMIDES.

Formula of Hydrobromic Acid HBr . Molecular weight 81.

Bromine: Source, Preparation, and Properties.—The acidulous radical of hydrobromic acid and other bromides is the element bromine, Br . (*Bromum*, B. P.). It occurs in nature chiefly as bromide of magnesium (MgBr_2) in sea-water and certain saline springs, and is commonly prepared from the *bittern*, or residual liquors of salt works. It may be liberated from its compounds by the process for chlorine from chlorides—that is, by heating with black oxide of manganese and sulphuric acid (*see* page 309). It is a dark-red volatile liquid, emitting an odour more irritating, if possible, than chlorine—of specific gravity 2.966, boiling point 154.4°F .

Quantivalence.—The atom of bromine, like that of chlorine, is univalent (Br'). The atomic weight of bromine is 80. Free bromine has the molecular formula Br_2 , bromide of bromine.

Fig. 38.



PREPARATION OF HYDROBROMIC ACID.

Hydrobromic Acid.—The bromide of hydrogen, hydrobromic acid, may be made by decomposing bromide of phosphorus by water— $\text{PBr}_5 + 4\text{H}_2\text{O} = 5\text{HBr} + \text{H}_3\text{PO}_4$. A small quantity is prepared by placing seven or eight drops of bromine at the

bottom of a test-tube, putting in fragments of glass to the height of about an inch or two, then ten or eleven grains of phosphorus, then another inch of glass, and finally a couple of inches of glass fragments slightly wetted with water, a delivery-tube being fitted on by a cork. The phosphorus combines readily, almost violently, with the bromine as soon as the vapour of the latter, aided by a little warmth from a flame, rises to the region of the phosphorus. The bromide of phosphorus thus formed then suffers decomposition by the water of the moist glass, phosphoric and phosphorous acids being produced. The hydrobromic acid gas passes over (heat being applied in the after part of the operation) and may be condensed in water or in solution of ammonia. The latter solution on evaporation yields bromide of ammonium.

Hager's method.—One hundred parts of sodium hyposulphite, 50 of bromine, and 10 of water are placed in a flask, and the generated gas is conducted into the upper portion of 140 parts of water contained in another vessel. When the gas begins to come over slowly, gentle heat is applied. The product is nearly 190 parts of liquid, containing 25 per cent, of real acid; sp. gr. 1.204. It should be kept in a cool dark place.

Squibb prefers to decompose solution of bromide of potassium by sulphuric acid, and, after removal of sulphate of potassium by crystallization, to distil the residual fluid. Wade prescribes an almost pure clear solution of the acid, made by shaking together 120 grains of bromide of potassium and 153 grains of crystallized tartaric acid in one ounce of water, and setting aside till precipitation of acid tartrate of potassium ceases.

Bromide of Potassium (KBr) is occasionally employed in pharmacy, and is the salt, therefore, which may be used in studying the reactions of this acidulous radical. The official method of making the salt has been alluded to under the salts of potassium (page 77).

Other bromides are seldom used; they may be prepared in the same way as, and closely resemble, the corresponding chlorides or iodides.

Bromide of Sodium crystallizes in anhydrous cubes (NaBr) from solutions at 110° F. or 120°, and in hydrated prisms (NaBr, 2H₂O) at ordinary temperatures.

Bromide of Ammonium (AmBr) (*Ammonii Bromidum*, B. P.) may be made by saturating hydrobromic acid with ammonia:

$\text{HBr} + \text{NH}_4\text{HO} = \text{NH}_4\text{Br} + \text{H}_2\text{O}$. It forms colourless crystals which become slightly yellow on exposure to air, is readily soluble in water, less so in spirit, and, when heated, sublimes.

Bromide or Iodide of Ammonium may also be made by mixing equivalent quantities of strong hot aqueous solutions of the corresponding potassium salts and of sulphate of ammonium. To the cooled liquids rectified spirit is added, which precipitates the sulphate of potassium. The spirit recovered by distillation of the clear liquid leaves the required salt as a residue in the retort.

Solution of Bromine, B. P., 10 minims in 5 ounces, is an aqueous solution, bromine being slightly soluble in water.

Hypobromites, *Bromates*, and *Perbromates*, analogous to hypochlorites, chlorates, and perchlorates, are producible.

Analytical Reactions (Tests).

First Analytical Reaction.—To a few drops of solution of a bromide (KBr , or NH_4Br) add solution of nitrate of silver; a yellowish-white precipitate of bromide of silver (AgBr) falls. Treat the precipitate successively with nitric acid and dilute ammonia, as described for the chloride of silver; it is only sparingly dissolved by the ammonia.

Second Analytical Reaction.—To solution of a bromide, add a drop or two of chlorine water, or a bubble or two of chlorine gas; then add a few drops of chloroform or ether, shake the mixture, and set the test-tube aside; the chlorine, from the greater strength of its affinities, liberates the bromine, which is dissolved by the chloroform or ether, the solution falling to the bottom of the tube in the case of the heavy chloroform, or rising to the top in the case of the light ether. Either solution has a distinct yellow, or reddish-yellow, or red colour, according to the amount of bromine present.

Notes.—This reaction serves for the isolation of bromine when mixed with many other substances. Excess of chlorine must be avoided, as colourless chloride of bromine is then formed. Iodides give a somewhat similar appearance; the absence of iodine must therefore be insured by a process given in the next section. The above solution in chloroform or ether may be removed from the tube by drawing up into a *pipette*

(small pipe—a narrow glass tube, usually having a bulb or expanded portion in the centre) the bromine fixed by the addition of a drop of solution of potash or soda, the chloroform or ether evaporated off, and the residue tested as described in the next reaction.

The above operation is frequently employed for synthetical purposes.

Third Analytical Reaction.—Liberate bromine from a bromide by the cautious addition of chlorine or chlorine-water, then add a few drops of cold decoction of starch; a yellow combination of bromine and starch, commonly termed "bromide of starch," is formed.

Decoction of starch is made by rubbing down two or three grains of starch with some drops of cold water, then adding much more water and boiling the mixture.

The above reaction may be varied by liberating the bromine by a little black oxide of manganese and a drop of sulphuric acid, the upper part of the inside of the test-tube being smeared over with some thick decoction of starch or thin starch-paste. Even sulphuric acid alone, if strong, liberates bromine from a bromide, the hydrogen of the hydrobromic acid first produced uniting with the oxygen of the sulphuric acid—the latter being reduced to sulphurous acid or even to hydrosulphuric acid.

HYDRIODIC ACID AND OTHER IODIDES.

Formula of Hydriodic Acid HI. Molecular weight 128.

Source.—The acidulous radical of hydriodic acid and other iodides is the element iodine (I). It occurs in nature chiefly as iodide of sodium and of magnesium in sea-water. Sea-weeds, sponges, and other marine organisms, which derive much of their nourishment from sea-water, store up iodides in their tissues; and it is from the ashes of these that supplies of iodine (*Iodum*, B. P.) are obtained.

Process.—The sea-weed ash or *kelp* is treated with water, insoluble matter thrown away, and the decanted liquid evaporated and set aside to allow of the deposition of most of the

sulphates, carbonates, and chlorides of sodium and potassium. The residual liquor is treated with excess of sulphuric acid, which causes evolution of carbonic and sulphurous or sulphuretted gases, deposition of sulphur and more sulphate of sodium, and formation of hydriodic acid. To the decanted liquid is added black oxide of manganese, and the mixture is then slowly distilled; the iodine sublimes, and is afterwards purified by resublimation.

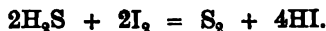


The analogy of chlorine, bromine, and iodine is well indicated by the fact that each is obtained from its compounds by the same reaction. Iodine is liberated from any iodide as bromine from bromides, or chlorine from chlorides—namely, by the action of black oxide of manganese and sulphuric acid.

Properties.—Iodine is a crystalline purplish-black substance; its vapour, readily seen on heating a fragment in a test-tube, is dark violet. Its vapours are irritating to the lungs; but a trace may be inhaled with safety (*Vapor Iodi*, B. P.). It melts at 239° F., boils at about 392°, and is entirely volatilized, the first portions containing any cyanide of iodine that may be present. The latter body occurs in slender colourless prisms, emitting a pungent odour.

Quantivalence.—The atom of iodine, like those of bromine and chlorine, is univalent* (I'). The atomic weight of iodine is 127; its molecular formula I_2 .

The Iodide of Hydrogen, or Hydriodic Acid, is a heavy, colourless gas. Its solution in water (*Acidum Hydriodicum Dilutum* of the United-States Pharmacopœia) may be made by passing sulphuretted hydrogen through water in which iodine is suspended.



* There is a compound of iodine having the formula ICl_3 . Iodine would at first sight therefore seem to be a trivalent element (I'''); and bromine and fluorine, from their close chemical analogy with iodine, would necessarily be regarded as trivalent also. From this aspect the position of chlorine would be anomalous. Probably, however, the compound is only a molecular combination of true chloride of iodine, ICl , with chlorine, Cl_2 . Iodine also forms, with iodide of potassium, a periodide or tri-iodide, KI_3 , which may be obtained in lustrous prismatic crystals. This, too, may have the formula $\text{KI} \cdot \text{I}_2$. A mercurio hexiodide (HgI_6 , perhaps $\text{HgI}_2 \cdot \text{I}_2$) is also known; and a periodide of ammonium $\text{NH}_4\text{I} \cdot \text{I}_2$.

Kolbe prepares hydriodic acid by adding to ten parts of iodine, contained in a flask in an atmosphere of carbonic acid gas, one part of phosphorus little by little, slowly. On the mixture of di- and tri-iodide of phosphorus is poured four parts of water. Abundance of hydriodic acid is evolved from this on the application of a gentle heat, and it is not contaminated by free iodine. Phosphoric acid remains.

Or iodine may be dissolved in bisulphide of carbon in a tall cylinder, water added, and sulphuretted hydrogen passed through the mixture. The water dissolves the hydriodic acid, the bisulphide retaining the separated sulphur. The aqueous solution only needs boiling for two or three minutes to remove excess of sulphuretted hydrogen.—*Winkler*.

Iodide of potassium (KI) is largely used in medicine, and hence is the most convenient iodide on which to experiment in studying the reactions of this acidulous radical. Solid iodine itself might be taken for the purpose; but its use and action in that state have already been alluded to in describing the iodides of potassium, cadmium, and mercury; its analytical reactions in the combined condition are those which may now occupy attention.

Solution of Iodine.—Iodine is slightly soluble in water (iodine-water), and readily soluble in an aqueous solution of iodide of potassium. Twenty grains of iodine and 30 of iodide of potassium, dissolved in 1 ounce of distilled water, form *Liquor Iodi*, B. P.; 32 grains of iodine and 32 of iodide of potassium, rubbed with 1 fluid drachm of proof spirit, and 2 ounces of lard gradually mixed in, form *Unguentum Iodi*, B. P. It is more soluble in spirit, or in a spirituous solution of iodide of potassium (*Tinctura Iodi*, B. P.). It combines with sulphur, forming an unstable greyish black solid iodide (S_2I_2), having a radiated crystalline structure (*Sulphuris Iodidum*, B. P., and *Unguentum Sulphuris Iodidi*). "If 100 grains be thoroughly boiled with water the iodine will pass off in vapour, and about 20 grains of sulphur remain."—*Brit. Pharm.*

Analytical Reactions (Tests).

First Analytical Reaction.—To a few drops of an aqueous solution of an iodide (e.g. KI) add solution of nitrate of silver; a light yellow precipitate of iodide of silver (AgI) falls. Pour away the supernatant liquid and treat the precipitate with

nitric acid, it is not dissolved; pour away the acid and then add dilute ammonia, it is only sparingly dissolved.

This reaction is useful in separating iodine from most other acidulous radicals, but does not distinguish iodine from bromine.

Ammonia, it will be remembered, dissolves chloride of silver readily; hence the presence of chloride of potassium in bromide or iodide may be detected by dissolving in water, adding excess of nitrate of silver, collecting the precipitate, washing, digesting in ammonia, filtering, and adding excess of nitric acid to the filtrate; a white curdy precipitate indicates a chloride (of potassium). Bromide and iodide of silver are, however, slightly soluble in ammonia. A better process is given on page 318.

Second Analytical Reaction.—Liberate iodine from an iodide by the cautious addition of chlorine, then add cold decoction of starch; a deep-blue combination of iodine and starch, commonly termed "iodide of starch," is formed.

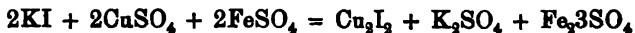
Starch is highly sensitive to the action of iodine; this reaction is consequently very delicate and characteristic. The reaction is not observed in hot liquids. Excess of chlorine must be avoided, or colourless chloride of iodine will be produced. Nitrous acid, or a nitrite acidulated with sulphuric acid, may be used instead of chlorine. Concentrated sulphuric acid also liberates iodine from iodides, the hydrogen of the hydriodic acid first produced uniting with the oxygen of the sulphuric acid—the latter (H_2SO_4) being reduced to sulphurous acid (H_2SO_3) or even to hydrosulphuric acid (H_2S).

In testing bromine for iodine the bromine must be nearly all converted into hydrobromic acid by dilute solution of sulphurous acid before the decoction of starch is added.

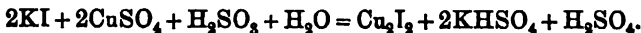
Ozone (O_3).—Papers soaked in mucilage of starch containing iodide of potassium form a test for free chlorine and nitrous acid, and are also employed by meteorologists to detect an allotropic or physically polymeric and energetic form of oxygen termed by Schönbein *ozone* (from $\delta\zeta\omega$, *ozō*, I smell). This substance liberates iodine from the iodide of potassium (with formation of iodide of starch), and is supposed to occur normally in the atmosphere, the salubrity or insalubrity of which is said to be dependent to some extent

on the presence or absence of ozone. The possible occurrence of nitrous or chlorinoid gases in the air, however, renders the test untrustworthy. Houzeau proposes to test for ozone by exposing litmus paper of a neutral tint soaked in a dilute solution of iodide of potassium: the potash set free by action of the ozone turns the paper blue. The same paper without iodide would indicate the extent to which the effect might be due to ammonia vapour. Ozone, or rather ozonized air, is produced artificially in large quantities on passing air through a box (Beane's Ozone-generator) highly charged with electricity. In the latter operation condensation of the volume of air, or, rather, of the oxygen in the air occurs. Small quantities are obtained by exposing in a loosely closed bottle a stick of phosphorus partially covered by water, but the product is mixed with peroxide of hydrogen. Ozone is a powerful bleaching, disinfecting, and general oxidizing agent; insoluble in water, soluble in oils of turpentine, cinnamon, and some other liquids. From experiments that have been made by Soret on the specific gravity of ozone, its molecular formula would seem to be O_3 , that of ordinary oxygen being O_2 . Its smell is peculiar.

Third Analytical Reaction.—To a neutral aqueous solution of an iodide add a solution containing one part of sulphate of copper to two and a half parts of green sulphate of iron, and well shake; a dirty white precipitate of cuprous iodide (Cu_2I_2) falls.



Or to the liquid containing an iodide add the solution of copper sulphate and some solution of sulphurous acid, and warm the mixture; cuprous iodide falls.



Separation of Chlorides, Bromides, and Iodides.—Chlorides and bromides are not affected in this way; the reaction is useful, therefore, in removing iodine from a solution in which chlorides and bromides have to be sought. The total removal of iodine by the former of the two modifications of the process is insured by supplementing the addition of the cupric and ferrous sulphates by a few drops of solution of potash or soda,

any acid which might be keeping cuprous iodide in solution being thereby neutralized, ferric or ferrous hydrate, precipitated at the same time, not affecting the reaction. Occasionally, too, it may be necessary to repeat the process with the filtrate before the last traces of iodine are removed. The second modification of the process is, on the whole, to be preferred.

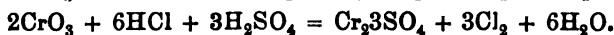
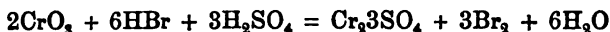
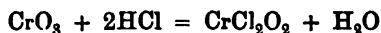
Chloride of the rare metal palladium performs a similar useful office in removing iodine, but not bromine or chlorine, from solutions.

Chlorides may be separated from bromides by taking advantage of the ready solubility of chloride of silver, and the slow and slight solubility of bromide of silver, in ammonia, especially in (a fair, not a great, excess of) ammonia containing chloride of silver.

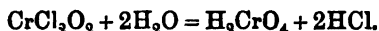
Siebold's test for the detection of chlorides when occurring with bromides is based upon the fact that a strong solution of permanganate of potassium liberates the bromine from moderately strong solutions of bromides containing a large excess of sulphuric acid. It may be performed in the following manner:—A strong solution of permanganate is added to the aqueous solution of bromide or iodide (containing not more than 1 in 40), strongly acidified by dilute sulphuric acid, until the permanganate ceases to be decolorized, and a copious precipitate of oxide of manganese is formed. The mixture is boiled for about five minutes to expel the bromine, and then filtered. The colourless filtrate is now quite free from bromine, and may be tested for chlorine by nitrate of silver. If a chloride be present, a small quantity of its chlorine is lost by this process; but the main portion always remains, provided that no undue excess of the permanganate be used. It is essential, therefore, that the filtrate should be colourless; for if it be coloured so as to indicate the presence of undecomposed permanganate, the loss of the greater part if not of the whole of the chlorine must be expected. If, on the other hand, too little permanganate be employed, a trace of bromide may be left in the filtrate. If the solution under examination should be very much stronger than 1 in 40, water should be added before boiling (just after the addition of the permanganate), in order to avoid a loss of HCl.

Chlorides may also be detected in bromides and iodides by taking advantage of the formation of chlorochromic anhydride (page 274) and the non-occurrence of corresponding compounds of bromine or iodine.

To a solution of a mixture of an iodide with a bromide and a chloride add a concentrated solution of sulphite of sodium, then a reagent prepared by mixing equal volumes of sulphuric acid and saturated solution of sulphate of copper, until no further precipitation of cuprous iodide occurs. Next add solution of soda to remove excess of sulphate of copper, filter and evaporate to dryness. Place the dried residue together with an equal bulk of red chromate of potassium in a dry test-tube fitted with a delivery tube, or into a small retort, and cover the mixture with sulphuric acid. Distil into water. Chromic anhydride and hydrochloric and hydrobromic acids are liberated by the sulphuric acid, and, reacting one upon another, form chlorochromic anhydride together with free bromine and chlorine.



The chlorochromic anhydride is decomposed by the excess of water into which it distils giving rise to chromic acid, which imparts its colour to the liquid, and hydrochloric acid, thus—



Chlorine gas escapes and the bromine is dissolved by the water. The coloured liquid is then shaken with chloroform, which removes the bromine, indicating bromides in the original substance. A yellow colour remaining is due to chromic acid, indicating chlorides in the original substance. Or add ammonia to the distillate—the colour due to bromine is thereby entirely removed, while that of the chromic compound is only slightly modified.

Fourth Analytical Reaction.—Iodides have been shown to be useful in testing for mercuric salts (*see* the Mercury reactions, p. 232); a mercuric salt (corrosive sublimate, for

example) may therefore be used in testing for iodides, a scarlet precipitate of mercuric iodide (HgI_2) being produced.

This reaction may be employed where large quantities of an iodide are present; but its usefulness in analysis is much impaired by the fact that the precipitate is soluble in excess of the dissolved iodide, or in excess of the mercuric reagent. Its colour and insolubility in water distinguish it from mercuric chloride, bromide, and cyanide, which are white soluble salts.

Fifth Analytical Reaction.—Iodides have also (*see* the Lead reactions, page 244) been shown to be useful in testing for lead salts; similarly a lead salt (acetate, for example) may be used in testing for iodides, a yellow precipitate of iodide of lead (PbI_2), soluble in hot water and crystallizing in yellow scales on cooling, being produced.

Chloride, bromide, and cyanide of lead are white; hence the above reaction may occasionally be useful in distinguishing iodine from the allied radicals. But iodide of lead is slightly soluble in cold water; hence small quantities of iodine cannot be detected by this reaction. (For *Iodates*, *see* Index.)

Analogies between Chlorine, Bromine, Iodine, and their Compounds.—These elements form a natural group or family, each distinct from the other, yet closely related. Moreover their dissimilarities are so curiously gradational as to irresistibly suggest the idea that some day we may find the differences between these bodies to be in degree rather than in kind. Thus chlorine is a gas and iodine a solid, while bromine occupies the intermediate condition. The atomic weight of bromine is nearly midway between those of chlorine and iodine. The same may be said of the weight of equal volumes of each in the gaseous state. The specific gravity of fluid chlorine is 1.33, of iodine 4.95, while bromine is nearly 3. Liquid chlorine is transparent, iodine opaque, bromine intermediate. The crystalline forms of the chloride, bromide, and iodide of a metal are commonly identical. One volume of either element in the gaseous state combines with an equal volume of hydrogen (at the same temperature) to form two volumes of a gaseous acid, very soluble in water (hydro-

chloric acid, hydrobromic acid, hydriodic acid). Many other analogies are traceable.

QUESTIONS AND EXERCISES.

417. State the method by which Bromine is obtained from its natural compounds.

418. Mention the properties of bromine.

419. How may the Bromides of Potassium and Ammonium be made?

420. By what reagents may bromides be distinguished from chlorides?

421. Whence is Iodine obtained?

422. By what process is iodine isolated?

423. State the properties of iodine.

424. What is the nature of Iodide of Sulphur?

425. Give the analytical reactions of iodides.

426. Which three substances may, indirectly, be detected by a mixture of iodide of potassium and mucilage of starch?

427. Describe a method by which iodides may be removed from a solution containing chlorides and bromides.

HYDROCYANIC ACID AND OTHER CYANIDES.

Formula of Hydrocyanic acid HCN or HCy .

Molecular weight 27.

History of Cyanogen.—The acidulous radical of hydrocyanic acid and other cyanides is a compound body, cyanogen (Cy). It is so named from *κύανος*, *kuanos*, blue, and *γεννάω*, *gennaō*, I generate, in allusion to its prominent chemical character of forming, with iron, the different varieties of Prussian blue. It was from Prussian blue that Scheele in 1782 first obtained what we now, from our knowledge of its composition, term hydrocyanic acid, but which he called Prussic acid. Cyanogen was isolated by Gay-Lussac in 1814, and was the first compound radical distinctly proved to exist.

Sources.—Cyanogen does not occur in nature, and is only formed from its elements under certain circumstances. It is found in small quantities among the gases of iron-furnaces,

and is produced to a slight extent in distilling coals for gas. In the form of ferrocyanide of potassium it is obtained abundantly by heating animal refuse containing nitrogen, such as the scrapings of horns, hoofs, and hides (5 parts), with carbonate of potassium (2 parts) and waste iron (filings, etc.) in a covered iron pot. The residual mass is boiled with water, the mixture filtered, and the filtrate evaporated and set aside for crystals to form. The cyanogen, produced from the carbon and nitrogen of the animal matter, unites with the potassium and afterwards, on boiling with water, with iron to form what is known as the yellow prussiate of potash (*Potassæ Prussias Flava*, B. P.), or ferrocyanide of potassium ($K_4Fe''Cy_6$), a compound occurring in four-sided tabular yellow crystals. It contains the elements of cyanogen, yet is not a cyanide, for it is not poisonous, and is otherwise different from cyanides: it will be further noticed subsequently. From this salt all cyanides are directly or indirectly prepared.

Cyanide of potassium (KCy), which is the most common, is procured by fusing eight parts of ferrocyanide of potassium with three of carbonate of potassium in a crucible; carbonic acid gas (CO_2) is evolved, iron (Fe) is set free, and cyanate of potassium (KCyO), a body that will be subsequently noticed, is formed at the same time:—



Double cyanides exist, such as the cyanide of sodium and silver ($NaCy$, $AgCy$) formed in the process (subsequently described) of quantitatively determining the amount of hydrocyanic acid in a liquid by a standard solution of nitrate of silver: these compounds have, more or less, the properties of their constituents. But other cyanogen compounds, not double cyanides, occur in which the cyanogen is so intimately united with a metal as to form a distinct radical: such are ferrocyanides and ferridcyanides—salts which will be noticed in due course.

Cyanogen, like chlorine, bromine, and iodine, is univalent (Cy). It may be isolated by simply heating mercuric cyanide ($HgCy_2$) or cyanide of silver ($AgCy$). It is a colourless gas, burning, when ignited, with a beautiful peachblossom-coloured flame.

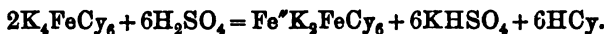
Mercuric cyanide is produced in crystals on dissolving 1 part of ferrocyanide of potassium in 15 parts of boiling water, adding 2 parts of mercuric sulphate, keeping the

whole hot for ten or fifteen minutes, and then filtering and setting aside to cool. In addition to mercuric cyanide (HgCy_2), mercury (Hg), ferric sulphate ($\text{Fe}_2\text{3SO}_4$), and sulphate of potassium (K_2SO_4) are formed. Any excess of ferrocyanide also gives Prussian blue by reaction with the ferric sulphate. A small flame of cyanogen may be obtained on heating a few crystals of mercuric cyanide in a short piece of glass tubing closed at one end, and applying a light to the other end as soon as evolution of gas commences: brown *paracyanogen* (C_3N_3) and mercury remain.

REACTIONS.

Diluted Hydrocyanic Acid.

Synthetical Reaction.—Dissolve 2 or 3 grains of ferrocyanide of potassium in 5 or 6 times its weight of water in a test-tube, add a few drops of sulphuric acid and boil the mixture, conveying the evolved gas by a bent glass tube (adapted to the test-tube by a cork) into another test-tube containing a little water; the product is a dilute solution of hydrocyanic acid. Made by this process in larger quantities and of a certain definite strength (2 per cent.), this solution is the *Acidum Hydrocyanicum Dilutum*, B. P. "A colourless liquid of peculiar odour. Specific gravity 0.997."



The details of the official process are as follows:—Dissolve $2\frac{1}{2}$ ounces of ferrocyanide of potassium in 10 ounces of water, add 1 fluid ounce of sulphuric acid previously diluted with 4 ounces of water and cooled. Put the solution into a flask or other suitable apparatus of glass or earthenware, to which are attached a condenser and a receiver arranged for distillation (*see* p. 138); and having put 8 ounces of distilled water into the receiver, and provided efficient means for keeping the condenser and receiver cold, apply heat to the flask, until by slow distillation the liquid in the receiver is increased to 17 fluid ounces.* Add to this 3 ounces of distilled

* This operation is peculiarly liable to those sudden and tumultuous evolutions of vapour, or "bumpings," or "*soubresauts*," which often interfere with successful distillation. Such phenomena occur, according to Tomlinson, whenever unaided heat has to overcome the great amount of adhesion naturally existing between certain liquids and vapours, or,

water, or as much as may be sufficient to bring the acid to the required strength, so that 100 grains (or 110 minims) of it, precipitated with a solution of nitrate of silver (*vide* paragraphs on quantitative analysis), shall yield 10 grains of dry cyanide of silver. The end of the condenser should pass quite into the receiver.

The residue of this reaction is acid sulphate of potassium (KHSO_4), which remains in solution, and ferrocyanide of potassium and iron ($\text{Fe}^+\text{K}_2\text{FeCy}_6$), an insoluble powder sometimes termed Everitt's yellow salt, from the name of the chemist who first made out the nature of the reaction. The latter compound becomes bluish green during the reaction, owing to absorption of oxygen.

Pure anhydrous hydrocyanic acid is a colourless, highly volatile, intensely poisonous liquid, solidifying when cooled to a low temperature.* It may be made by passing sulphuretted hydrogen over mercuric cyanide. The official solution of the acid is fairly stable, but is said to be rendered more so by the presence of a minute trace of sulphuric or hydrochloric acid. A stronger acid is liable to assimilate the elements of water, and yield formate of ammonium (NH_4CHO_2). Solu-

rather, between the normal liquid and those particles of it which, becoming strongly heated at the heated part of the vessel, have assumed the condition of particles of dissolved vapour, and which would at once pass from this condition into that of permanent vapour but for adhesion. Ordinarily a glass or other surface is not absolutely clean, but is more or less covered with specks, traces of materials deposited from the air, the fingers, cloths, etc. Some liquids seem to have little or no adhesion for these materials, while certain vapours have greater adhesion for the films than for the liquids. Hence, in ordinary regular ebullition the vapours accumulate on the films, and then at once become subject to the pressure of the mass of fluid, and so pass off in bubbles. But when the films are absent, or have become removed during distillation, the heat accumulates until it is sufficient to overcome the adhesion of the superheated particles, and these are then, all of them at once, converted into vapour, the liquid commonly boiling over, sometimes even bursting the vessel. "Bumping" would be prevented by the introduction of fragments of substances for which vapour-particles have adhesion, but no known substance has this property in an absolute degree. Fragments of tobacco-pipe or pumice-stone, pieces of cork, thick paper, resin, sulphur, platinum wire, etc., are all useful when there is no chemical action between them and the liquid. Mr. Tomlinson very strongly recommends cocoanut-shell charcoal to be used whenever practicable. A slow current of gas, such as hydrogen, air, or carbonic acid gas, also usefully promotes escape of vapour from a liquid. A jet of steam prevents it, but is not always applicable.

* Traces are formed when electricity passes between carbon poles in slightly moist air (Dewar).

tions of hydrocyanic acid often become brown by formation of what is, apparently, *paracyanogen* (C_3N_3). According to Williams aqueous hydrocyanic acid containing 20 per cent. of glycerine can be kept for an almost indefinite length of time.

Note.—A few drops of diluted hydrocyanic acid so placed that its vapour may be inhaled, forms the *Vapor Acidi Hydrocyanici*, B. P., or Inhalation of Hydrocyanic Acid.

Hydrocyanic acid also occurs in cherry-laurel water and bitter-almond water (*vide* Index).

The methods of determining the strength of solutions of hydrocyanic acid will be described in connection with volumetric and gravimetric quantitative analysis. They are based on the formation of cyanide of silver, and its solubility in solution of cyanide of potassium, as described in the next reaction.

The Hydrocyanic Acid used in pharmacy is extremely liable to variation in strength. It should frequently be tested volumetrically.

Analytical Reactions (Tests).

First Analytical Reaction.—To a few drops of the hydrocyanic acid solution produced in the above reaction, or to any solution of a cyanide, add excess of solution of nitrate of silver; a white precipitate of cyanide of silver ($AgCy$) falls. When the precipitate has subsided pour away the supernatant liquid and place half of the residue in another test-tube: to one portion add nitric acid, and notice that the precipitate does not dissolve; to the other add ammonia, and observe that the precipitate is insoluble, or only sparingly soluble. (Chloride of silver, which is also white, is readily soluble in ammonia.) Cyanide of silver dissolves in solution of cyanides of alkali-metals, soluble double cyanides being formed (*e.g.* KCy , $AgCy$).

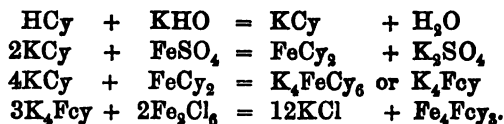
Solubility of precipitates in strong solutions of salts.—Cyanide of silver and many other precipitates insoluble in acids (similar remarks apply to precipitates insoluble in alkalies) are often soluble in the strong saline liquids formed by the addition of acids and alkalies to each other. Hence the precaution of adding the latter reagents to separate portions of a precipitate,

or of not adding the one until the other has been poured away.

Cyanogen in an insoluble cyanide, such as cyanide of silver itself, is readily recognised on heating the substance in a short piece of glass tubing closed at one end like a test-tube and drawn out at the other end, so as to have but a small opening; on applying a flame, the escaping cyanogen ignites and burns with a characteristic peach-blossom tint.

Antidote.

Second Analytical Reaction.—To a dilute solution of hydrocyanic acid, or a soluble cyanide, add a few drops of solution of a ferrous salt and a drop or two of solution of a ferric salt (ferrous sulphate and ferric chloride are usually at hand); to the mixture add potash, soda, magnesia, or carbonate of sodium, and then hydrochloric acid; a precipitate of Prussian blue remains. The decompositions may be traced in the following equations:—



The test depends on the conversion of the cyanogen into ferrocyanogen by aid of the iron of a ferrous salt, and the combination of the ferrocyanogen, so produced, with the iron of a ferric salt.

Hence a mixture of green sulphate of iron, solution of perchloride of iron, and either magnesia or carbonate of sodium is the recognised *antidote* in cases of poisoning by hydrocyanic acid or cyanide of potassium.

In such an alkaline mixture the poisonous cyanide, by reaction with ferrous hydrate, is at once converted into innocuous ferrocyanide of potassium or sodium, etc.: should the mixture become acid, the ferric salt present reacts with the soluble ferrocyanide forming insoluble Prussian blue, which is also inert. From the rapidity of the action of these poisons, however, there is seldom time to prepare an antidote. Emetics, the stomach-pump, the application of a stream of cold water to the spine, and the above antidote form the usual treatment.

Third Analytical Reaction.—To solution of hydrocyanic acid add ammonia and common yellow sulphhydrate of ammonium, and evaporate the liquid nearly or quite to dryness in a small dish, occasionally adding ammonia till the excess of sulphhydrate of ammonium is decomposed; add water and acidify the liquid with hydrochloric acid, and then add a drop of solution of a ferric salt; a blood-red solution of sulphocyanate of iron will be formed.

This is a very delicate reaction. Some free sulphur in the yellow sulphhydrate of ammonium unites with the alkaline cyanide and forms sulphocyanate ($2\text{AmCy} + \text{S}_2 = 2\text{AmCyS}$); the ammonia combines with excess of free sulphur and forms, among other salts, sulphhydrate of ammonium, the whole of which is removed by the ebullition. If the liquid has not been evaporated far enough, sulphhydrate of ammonium may still be present, and give black sulphide of iron on the addition of the ferric salt.

Hydrocyanic acid in the blood.—According to Buchner the blood of animals poisoned by hydrocyanic acid, instead of coagulating as usual, remains liquid and of a clear cherry-red colour for several days. In one case he obtained the reactions of the acid on diluting and distilling the blood fifteen days after death, and applying the usual reagents to the distillate. Aqueous solution of peroxide of hydrogen changes such blood to a deep brown colour.

Schönbein's test for hydrocyanic acid is said to be extremely delicate. Filtering-paper is soaked in a solution of 3 parts of guaiacum resin in 100 of alcohol. A strip of this paper is dipped in a solution of 1 part of sulphate of copper in 50 of water; a little of the suspected solution is placed on this paper and exposed to the air, when it immediately turns blue. Or the paper may be placed over the neck of an open bottle of medicine supposed to contain hydrocyanic acid, or otherwise exposed to the vapour of the acid.

QUESTIONS AND EXERCISES.

428. Write a paragraph on the history of cyanogen.
429. Mention the source of the cyanogen of cyanides.
430. How is Ferrocyanide of Potassium prepared?

431. What is the formula of ferrocyanide of potassium ?
432. Is ferrocyanide of potassium poisonous ?
433. Write an equation expressive of the reaction which ensues when ferrocyanide and carbonate of potassium are brought together at a high temperature.
434. What are the properties of cyanogen ? How may it be obtained in a pure condition ?
435. How is mercuric cyanide prepared ?
436. How much real hydrocyanic acid is contained in the official liquid ?
437. Give details of the preparation of hydrocyanic acid, and an equation of the reaction.
438. State the proportion of water that must be added to an aqueous solution containing 15 per cent. of hydrocyanic acid to reduce the strength to 2 per cent.—*Ans.* $6\frac{1}{2}$ to 1.
439. What are the characters of pure undiluted hydrocyanic acid ? How may it be obtained ?
440. Enumerate the tests for cyanogen, giving equations.
441. Explain the action of the best antidote in cases of poisoning by hydrocyanic acid or cyanide of potassium.
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NITRIC ACID AND OTHER NITRATES.

Formula of Nitric Acid HNO_3 . Molecular weight 63.

Introduction.—The group of elements represented by the formula NO_3 is that characteristic of nitric acid and all other nitrates ; hence it is expedient to regard these elements as forming an acidulous radical, which may be termed *the nitric radical*. Like the hypothetical basylous radical ammonium (NH_4), this supposed acidulous radical (NO_3) has not been isolated. Possibly it is liberated when chlorine is brought into contact with nitrate of silver ; but if so, its decomposition into white crystalline nitric anhydride (N_2O_5) and oxygen (O) is too rapid to admit of its identification.

Sources.—The nitrogen and oxygen of the air combine and ultimately form nitric acid whenever a current of electricity (as in the occurrence of lightning) passes. The nitrates found in rain may partly or wholly thus originate. The oxidation of ammoniacal matter, and of the nitrogenous constituents of animal and vegetable matters in the soil, favoured, perhaps,

by the darkness, by the occurrence of carbonate of calcium, and, possibly, by the presence of some low form of vegetable life acting as a ferment, result in the production of nitrates. Hence nitrates are commonly met with in waters, soils, and the juices of plants. In the concentrated plant-juices termed medicinal "Extracts," small prismatic crystals of nitrate of potassium may occasionally be observed. (The cubical crystals often met with on extracts are chloride of potassium.) Nitric acid and other nitrates are obtained from nitrates of potassium and sodium, and these from the surface layers of the soil of tropical countries. *Nitrate of potassium* or *prismatic nitre* (from the form of its crystals) is chiefly produced in and about the villages of India. The natives simply scrape the surface of waste grounds, mud heaps, banks, and other spots where a slight incrustation indicates the presence of appreciable quantities of nitre, mix the scrapings with wood-ashes (carbonate of potassium, to decompose the nitrate of calcium always present), digest the mixture in water, and evaporate the liquor. The immediate product is purified by careful recrystallizations, and is sent into commerce in the form of white crystalline masses or fragments of striated six-sided prisms. Besides its use in medicine (*Potassæ Nitras*, B. P.), it is employed in very large quantities in the manufacture of gunpowder. *Nitrate of Sodium* (*Sodæ Nitras*, B. P.) occurs in deposits from 3 inches to 3 yards in thickness on and near the surface, and at any depth down to about 30 feet, in many parts of Peru, Bolivia, and Chili, but more especially in the district of Atacama. The mineral is termed *caliche*, and commonly contains 50 per cent. of nitrate of sodium. The latter is distinguished as Chili saltpetre or (from the form of its crystals—obtuse rhomboids, not true cubes) cubic nitre, and is chiefly used as a manure and as a source of nitric acid, its tendency to absorb moisture unfitting it for use in gunpowder. In many parts of Europe nitrate of potassium is made artificially by exposing heaps of animal manure, refuse, ashes, and soil to the action of the air and the heat of the sun; in the course of a year or two the nitrogen of the animal matter becomes oxidized to nitrates; the latter are removed by washing.

Note.—The word *nitric* is from *nitre*, the English equivalent of the Greek *νίτρον* (*nitron*), a name applied to certain natural deposits of *natron* (carbonate of sodium), for which nitrate of potassium seems at first to have been mistaken. *Saltpetre* is

simply *sal petrae*, salt of the rock, in allusion to the natural origin of nitrate of potassium. *Sal prunella* (from *sal*, a salt, and *pruna*, a live coal) is nitrate of potassium melted over a fire and cast into cakes or bullets.

The nitric radical is univalent (NO_3').

Constitution of Salts.

It is here necessary again to caution the reader against regarding salts as invariably possessing a known constitution, or supposing that they always possess two or more sides, or contain definite radicals. The erroneous conception which, of all others, is most likely to be imperceptibly formed is that of considering salts to be binary bodies. For, first, the names of salts are necessarily binary. A student hears the names "sulphate of iron," "sulphate of copper," and simultaneously receives the impression that each salt has two sides, copper or iron occupying one, and something indicated by the words "sulphate of" the other. Such words as "vitriol," green or blue, or "nitre," would perhaps implant unitary ideas in the mind; but it is simply impossible to give such names to all salts as will convey the impression that each salt is a whole, and therefore unitary. The name "sulphate of potash" produces binary impressions; and the less incorrect name "sulphate of potassium" is in this respect no better. Secondly, it is impracticable to study salts as a whole. Teachers are unanimous in the opinion that students should first master the reactions characteristic of the metals in salts, and then the residues which, with those metals, make up the salts, or *vice versa*. It is not only impracticable, but impossible, to study salts as a whole; binary ideas concerning them are therefore almost inevitably imbibed. We come to regard a salt as a body which splits up in one direction only, look upon nitre, for instance, and all other nitrates, as containing NO_3 and a metal, M ; whereas KNO_3 may be split up into KNO_2 and O ; or into K_2O , N_2 , and O_5 ; or may contain K_2O and N_2O_5 . These are the chief disadvantages attending the employment of the binary hypothesis in studying chemical compounds: if they be borne in mind, the hypothesis may be freely used without much danger of permanent mental bias. Thus in nitre let the group of elements (NO_3) which, with potassium, makes up the whole salt be called the nitric radical, the name of the latter being directly derived from its hydrogen

salt. Similarly allow the acidulous residues of other salts of metals to be termed respectively the chloric, acetic, sulphurous, sulphuric, carbonic, oxalic, tartaric, phosphoric, citric, boracic radicals. In short, these compound radicals should be regarded as groupings common to many salts, and which may usually be transferred without any apparent breaking or splitting; at the same time we must be prepared to find that occasionally a salt divides in other directions. In this way perhaps erroneous impressions will gain least hold on the mind, and a way be left open for the easy entrance of new truths, should the real constitution of salts be discovered.

Formerly salts (such as sulphate of magnesium) were regarded as containing (a) an oxide of a metal (MgO) and an anhydride (SO_3), the latter being incorrectly called an acid (sulphuric acid); or (b) as containing two simple radicals (e.g. KI , $NaCl$, KCy , HgS)—the former being called *oxyacid salts* or *oxysalts*, and the latter *haloid salts* (from *ἄλς*, *als*, seasalt, and *εἶδος*, *eidos*, likeness). Such distinction is no longer maintained, the two classes being merged. This is an important educational gain on the side of simplicity; for whereas under the old system much time was necessarily expended before salts of a metal and salts of the oxide of that metal could be distinguished (e.g. KI and K_2O , SO_3), now, all salts being regarded as salts of the metals themselves (e.g. KI and K_2SO_4), no such distinction is necessary.

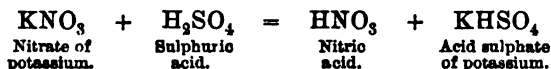
REACTIONS.

Nitric Acid.

Synthetical Reaction.—To a fragment of nitrate of potassium or nitrate of sodium in a test-tube add a drop or two of sulphuric acid, and warm; nitric acid (HNO_3) is evolved in vapour. The fumes may be condensed by a bent tube fitted to the test-tube, not by a cork as for hydrochloric acid—because the nitric vapours would strongly act on it—but by plaster of Paris, a paste of which sets hard on being set aside for a short time, and is unaffected by the acid.

On a somewhat larger scale nitric acid may be prepared by heating, in a stoppered or plain retort, a mixture of equal weights of nitrate of potassium and sulphuric acid;

the acid distils over, and acid sulphate of potassium remains behind:—



Half the quantity of sulphuric acid may be taken; but in that case neutral sulphate of potassium (K_2SO_4) is produced, which, from its hard, slightly soluble character, is removed with difficulty from the retort. On the manufacturing scale the less proportion is used; but instead of retorts iron cylinders are employed, from which the residual salt is removed by chisels. Moreover the cheaper sodium salt is the nitrate from which manufacturers usually prepare nitric acid, seven parts of nitrate of sodium and four of sulphuric acid being employed.

Note.—The acid sulphate of potassium is readily converted into neutral sulphate (*Potassæ Sulphas*, B. P.) by dissolving in water, adding carbonate of potassium until effervescence ceases to occur, filtering, and setting aside to crystallize.

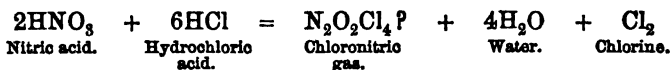
Pure nitric acid (HNO_3) is a colourless liquid, somewhat difficult of preparation; its specific gravity is 1.52. The strongest acid met with in commerce has a sp. gr. of 1.5, and contains 98 per cent. of real nitric acid (HNO_3); it fumes disagreeably, is unstable, and, except as an escharotic, is seldom used. The British Pharmacopœia contains two acids:—*Acidum Nitricum*, prepared as above, of sp. gr. 1.42, and containing 70 per cent. of real acid (HNO_3); and another, *Acidum Nitricum Dilutum*, sp. gr. 1.101, containing nearly $17\frac{1}{4}$ (17.44) per cent. Either of the stronger liquids, although containing water, is usually simply termed “nitric acid.” The official nitric acid, of sp. gr. 1.42, is a definite hydrous acid ($2\text{HNO}_3, 3\text{H}_2\text{O}$); it distils at 250°F . without change. If a weaker acid be heated it loses water, if a stronger acid be heated it loses nitric acid, until the density of 1.42 is reached. *Aqua fortis* is an old name for nitric acid (*Aqua fortis simplex*, sp. gr. 1.22 to 1.25; *Aqua fortis duplex*, 1.36). The strength of a specimen of nitric acid is determined by volumetric analysis. *Nitric anhydride* (N_2O_5), sometimes, but erroneously, called *anhydrous nitric acid*, is a solid crystalline substance formed on passing dry chlorine over dry nitrate of silver.

Metals reduce nitric acid to nitrous acid and to the various oxides of nitrogen or even to nitrogen itself, according to the

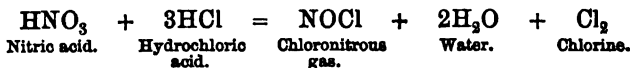
strength of acid, temperature, and amount of nitrate present. Not unfrequently nitrate of ammonium is simultaneously formed. Thus, with zinc,



Aqua Regia.—Three fluid ounces of nitric acid (B. P.), four of hydrochloric acid (B. P.), and twenty-five of water form the *Acidum Nitrohydrochloricum Dilutum* of the British Pharmacopoeia. The acids are ordered to be mixed twenty-four hours before dilution to insure mutual decomposition and full development of the chief active product, chlorine:—



In the later stages of the reaction, the decomposition expressed in the following equation also probably occurs:—

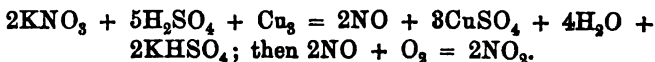


The same reaction occurs if the acids are mixed after dilution, but is not complete for a week or a fortnight (Tilden). The undiluted mixture of acids is known as *aqua regia*, so called from its property of dissolving gold, the “king of metals.”

This “diluted nitrohydrochloric acid” is quite strong enough to attack organic matter with evolution of nitrous gases, hence should not be dispensed with tinctures, etc., in too concentrated a form.

Analytical Reactions (Tests).

First Analytical Reaction.—To a solution of any nitrate (e.g. KNO_3) add sulphuric acid, and then copper turnings, and warm; colourless nitric oxide gas (NO) is evolved, which at once unites with the oxygen in the tube, giving *red fumes* of nitric peroxide or peroxide of nitrogen (NO_2).



Performed on a larger scale, in a vessel to which a delivery-tube is attached, the reaction of nitric acid on copper becomes

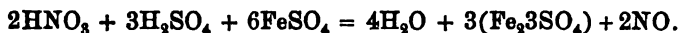
of synthetical interest, being the process for the preparation of nitric oxide gas for the purposes of chemical experiment.

Small amounts of a nitrate may be overlooked by this test, the colour of the red fumes not being very intense.

Undiluted nitric acid poured on to copper turnings gives dense red vapours of *nitrous acid* (HNO_2), *nitrous anhydride* (N_2O_3), *nitric peroxide* (NO_2), *nitric oxide* (NO), and even *nitrogen* (N_2), the reaction varying somewhat according to the temperature of the mixture and (Ackworth) the amount of nitrate of copper in solution. Diluted nitric acid gives nitric oxide, $\text{Cu}_3 + 8\text{HNO}_3 = 3(\text{Cu}_2\text{NO}_3) + 4\text{H}_2\text{O} + 2\text{NO}$.

Second Analytical Reaction.—To a cold solution of a nitrate, even if very dilute, add three or four crystals of sulphate of iron, shake gently for a minute in order that some of the sulphate may become dissolved, and then pour eight or ten drops of strong sulphuric acid down the side of the test-tube, so that it may form a layer at the bottom of the vessel; a reddish purple or black coloration will appear between the acid and the supernatant liquid.

This is a very delicate test for the presence of nitrates. The black colour is due to a solution or, perhaps, combination of nitric oxide with a portion of the ferrous salt. The nitric oxide is liberated from the nitrate by the reducing action of the hydrogen of the sulphuric acid, the sulphuric radical of which is absorbed by the ferrous sulphate, the latter salt becoming ferric sulphate.



The process of oxidation is one frequently employed in experimental chemistry; and nitrates, from their richness in oxygen, but more especially because always at hand, are the oxidizers usually selected for the purpose. In the operation they generally split up in one way, namely into oxide of their basylous radical nitric oxide gas and available oxygen. Thus hydrogen nitrate (nitric acid) yields oxide of hydrogen (water) and the other bodies mentioned, as shown in the following equation:—



When nitrates, other than nitric acid, are used for the purpose of oxidation, a stronger acid, generally sulphuric, is commonly

added in order that nitric acid may be formed, the hydrogen nitrate splitting up more readily than other nitrates.

The five oxides of nitrogen have now been mentioned, namely :—

Nitrous oxide	N_2O	} or {	N_2O
Nitric oxide*	NO		N_2O_2
Nitrous anhydride	N_2O_3		N_2O_3
Nitric peroxide*	NO_2		N_2O_4
Nitric anhydride	N_2O_5		N_2O_5

Nitrous oxide is a colourless gas, not altered on exposure to air; nitric oxide is also colourless, but gives red fumes in the air; nitrous anhydride is a red vapour condensible to a blue liquid; nitric peroxide is a red vapour condensible to an orange liquid; nitric anhydride is a colourless crystalline solid. The two anhydrides by absorbing water yield respectively nitrous acid (HNO_2) and nitric acid (HNO_3). Nitrous oxide is also probably an anhydride, the acid of which would doubtless have the formula HNO , while the silver and sodium salts certainly have the formulæ $AgNO$ and $NaNO$, $3H_2O$ (Divers; Menke). The above series of compounds forms a good illustration of the doctrine of multiple proportions (p. 45).

Third Analytical Reaction.—Direct the blowpipe-flame on to charcoal until a spot is red-hot; now place on the spot a fragment of a nitrate; deflagration ensues.

This reaction does not distinguish nitrates from chlorates. It is insufficient for the recognition of very small quantities of either class of salts, especially when they are mixed with other substances.

Gunpowder is an intimate mechanical mixture of 75 parts of nitre, 15 to $12\frac{1}{2}$ parts of charcoal, and 10 to $12\frac{1}{2}$ parts of sulphur. In burning it may be said to give sulphide of potassium (the white smoke, K_2S), nitrogen (N), carbonic oxide (CO), and carbonic acid (CO_2) gases, though the decomposition is seldom complete. The sudden production of a large quantity of heated gas from a small quantity of a cold solid is sufficient to account for all the effects of gunpowder.

* The specific gravities of these gases indicate that NO and NO_2 are the correct formulæ, and not N_2O_2 and N_2O_4 .

Fourth Analytical Reaction.—To nitric acid or other nitrate add solution of “sulphate of indigo;” the colour is discharged.

“*Solution of Sulphate of Indigo*,” B. P. (Sulphindyllic or Sulphindigotic Acid), is made by digesting 5 grains of dry finely powdered indigo in a small quantity of strong sulphuric acid in a test-tube for an hour, the mixture being kept hot by a water-bath; the blue liquid is then poured into 10 ounces of sulphuric acid, the whole well shaken, set aside, and the clear liquid decanted. Free chlorine also destroys the colour of this reagent.

Indigo, B. P. (C_8H_5NO), is a blue colouring-matter deposited when infusion of various species of *Indigofera* is exposed to air and slight warmth. Under these circumstances, *indican*, a yellow transparent amorphous substance, soluble in water, breaks up into indigo, which is insoluble and falls as a sediment, and a sort of sugar termed *indiglucin*. The indigo is collected, drained, pressed, and dried. By action of deoxidizing agents indigo is converted into soluble colourless *indigogen*, *reduced indigo*, or *white indigo*: 1 part of powdered indigo, 2 of green sulphate of iron, 3 of slaked lime, and 200 of water, shaken together and set aside in a well-closed bottle, gives this *colourless indigo*. A piece of yarn, calico, or similar fabric, dipped into such a solution and exposed to air, becomes dyed blue, deposition of insoluble indigo-blue occurring within the cells and vessels of the fibre. This operation is readily performed on the small scale, and forms a good illustration of the characteristic feature of the art of dyeing—namely, the introduction of soluble colouring-matter into a fabric by permeation of the walls of its cellular and vascular tissue, and the imprisonment of that colouring-matter within the cells and vessels by conversion into a solid and insoluble form (*vide* also p. 152).

Pure indigo, or *indigotin*, may be obtained in beautiful needles by spreading a paste of indigo and plaster of Paris on a tin plate, and when dry placing a lamp underneath, moving the latter from place to place as the indigo sublimes and condenses on the surface of the plaster. It may also be obtained in crystals by gently boiling finely powdered indigo with aniline, filtering while hot, and setting aside; these crystals may be washed with alcohol. Hot paraffin may be employed instead of aniline. Indigo is now produced] artificially.

Toluene, from coal tar, is, by Perkin's process, converted into cinnamic acid, this into a nitroderivative, and this again into orthonitropropionic acid. From the latter, alkali and grape sugar deposit crystalline indigo (Baeyer).

Distinction between nitric acid and other nitrates.—Presence of the nitric radical in a solution having been proved by the above reactions, its occurrence as the nitrate of a metal is demonstrated by the neutral, or nearly neutral, deportment of the liquid with test-paper and the detection of the metal—its occurrence as nitric acid by the sourness of the liquid to the taste and the effervescence produced on the addition of a carbonate.

Antidote.—In cases of poisoning by strong nitric acid, solution of carbonate of sodium (common washing soda) or magnesia and water may be administered as antidotes.

QUESTIONS AND EXERCISES.

442. Trace the origin of nitrates.
 443. In what does cubic nitre differ from prismatic nitre?
 444. Describe a process by which nitrate of potassium may be obtained artificially.
 445. State the difference between nitrate of potassium, nitre, saltpetre, and sal prunella.
 446. What group of elements is characteristic of all nitrates? and what claim has this group to the title of radical?
 447. Mention the usual theory regarding the manner in which atoms are arranged in reference to each other in such salts as nitrate of potassium.
 448. How is the official Nitric Acid prepared?
 449. Give the properties of nitric acid.
 450. What reactions occur when strong nitric and hydrochloric acids are mixed?
 451. How is nitric oxide prepared?
 452. Enumerate and explain the tests for nitrates.
 453. Into what substances does nitric acid usually split when employed as an oxidizing agent?
 454. How is nitrous oxide prepared?
 455. Enumerate the five oxides of nitrogen.
 456. What is the nature of gunpowder?
-

457. Write a few sentences on the chemistry of indigo, one of the tests for nitric acid.

458. How is nitric acid distinguished from other nitrates?

459. What quantity of cubic nitre will be required to produce ten carboys of official nitric acid, each containing 114 pounds?—
Ans. 1076 $\frac{1}{4}$ pounds.

CHLORIC ACID AND OTHER CHLORATES.

Formula of Chloric Acid HClO_3 . Molecular weight 84.5.

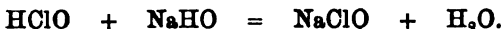
Chlorates are made from hypochlorites.

Hypochlorous Acid (HClO) and other Hypochlorites.

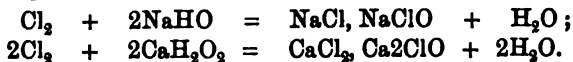
Place a few grains of red oxide of mercury in a test-tube, half-fill the tube with chlorine water and well shake the mixture; the resulting liquid is a solution of hypochlorous acid, mercuric oxychloride remaining undissolved:—



By the metathesis (double decomposition) of hypochlorous acid and oxides or hydrates, other pure hypochlorites are formed:—



The direct action of chlorine on metallic hydrates and some carbonates is supposed to give a compound of chloride and hypochlorite, as described in connection with the synthetical reactions of Sodium (page 90, *Liquor Sodæ Chloratæ*, B. P.) and, especially, calcium (page 122, *Calx chlorata*, B. P.).



The condition of the chlorine in these bodies is not satisfactorily made out; so that their constitution is not definitively determined. The action of acids on them results in the evolution of chlorine; hence the great value of the calcium compound (chlorinated lime, or chloride of lime) in bleaching-operations:—

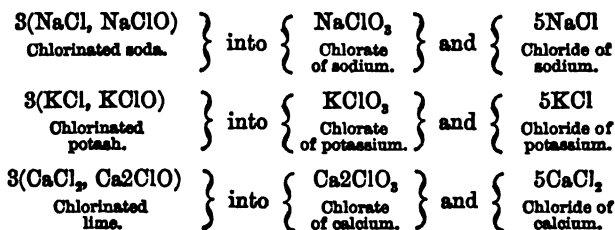


The solubility of hypochlorites in water, their peculiar odour, greatly intensified on the addition of acid, and their bleaching-powers (*see* the above calcium reaction) are the characters on which to rely in searching for hypochlorites.

Chlorates.

The group of elementary atoms represented by the formula ClO_3 is that characteristic of chloric acid and all other chlorates; hence it is expedient to regard it as being an acidulous radical, which may be termed the chloric radical. Like the nitric radical, it has not been isolated. Chloric anhydride (Cl_2O_5), unlike nitric anhydride, has not yet been obtained in the free condition.

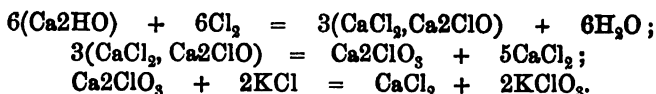
Chlorates are artificial salts. They are formed by simply boiling aqueous solutions of the common bleaching salts (chlorinated lime, chlorinated soda, chlorinated potash). Heat thus converts



One chlorate may also be made from another by double decomposition. In making chlorates economically the chlorinated salt is, of course, at once converted into chlorate.

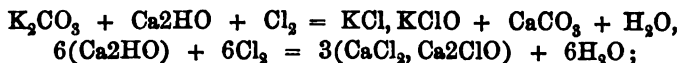
Chlorate of Potassium.

Thus Chlorate of Potassium (*Potassæ Chloras*, B. P.) is commercially made by saturating with chlorine gas a moistened mixture of 3 parts of chloride of potassium and 10 of slaked lime, and well boiling the product. Chlorinated lime is first formed; this, on continued boiling with water, splits up into chloride of calcium and chlorate of calcium; and the latter reacting on the chloride of potassium yields chloride of calcium and chlorate of potassium.

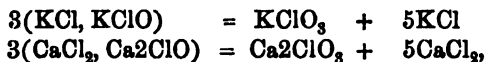


The operation may be conducted on a small scale by rubbing together in a mortar the above proportions of ingredients in ounces or half-ounces, adding enough water to make the whole assume the character of damp lumps, placing the porous mass in a funnel (loosely plugged with stones or pieces of glass) and passing chlorine gas (p. 18) through the mass by attaching the tube delivering the gas to the neck of the funnel. When the whole mass has become of a slight pink tint (due to a trace of permanganate) it should be turned into a dish, *well* boiled with water, filtered, the filtrate evaporated if necessary, and set aside; the chlorate of potassium crystallizes out in colourless rhomboidal plates, chloride of calcium remaining in the mother-liquor.

In the official process, carbonate of potassium instead of chloride is used; but otherwise it is similar to the method just described. Chlorinated potash and chlorinated lime are first formed—



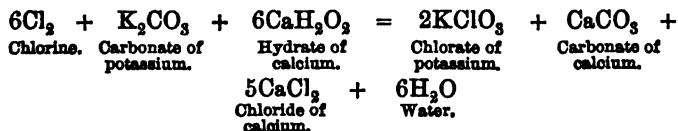
these on boiling with water split up into chlorates and chlorides—



the whole of the chloride of potassium and chlorate of calcium finally yielding chlorate of potassium and chloride of calcium,

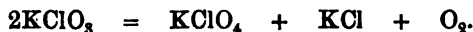


Neglecting intermediate decompositions, the reactions may be represented by the following equation :—



Chlorate of potassium is soluble in water to the extent of 6 or 7 parts in 100 at common temperatures. It is usually administered medicinally in aqueous solution, sometimes also in lozenges (*Trochisci Potassæ Chloratis*, B. P.). Chlorate of potassium must, on no account, be rubbed with sulphur in a mortar or otherwise, friction of such a mixture resulting in violent explosion.

Chlorate of potassium, when heated, yields chloride of potassium and oxygen, and is the salt commonly employed in the preparation of the gas for experimental purposes. But if the action be arrested when one-third of the oxygen has escaped, the residual salt is found to contain perchlorate of potassium (KClO_4):—



Perchloric Acid (HClO_4).—Crude perchlorate of potassium, obtained as just indicated, is boiled (in a fume-cupboard) with hydrochloric acid to decompose any chlorate that may be remaining, and then separated from chloride by washing and crystallization, chloride being far more soluble in water than the perchlorate. Perchloric acid is then obtained by distilling the perchlorate of potassium with sulphuric acid; it is quite stable, and is occasionally administered in medicine.

Chloric Acid (HClO_3) may be isolated, but is unstable, quickly decomposing into chlorine, oxygen, and perchloric acid; some other chlorate (e.g. KClO_3) must therefore be used in studying the reactions of the chloric radical.

Table of the Chlorine Acids.

Hydrochloric acid	HCl .
Hypochlorous acid	HClO .
Chlorous acid	HClO_2 .
Chloric acid	HClO_3 .
Perchloric acid	HClO_4 .

The chloric radical is univalent (ClO_3'). The acidulous radicals of the other chlorine acids are also univalent, as indicated in the foregoing formulæ.

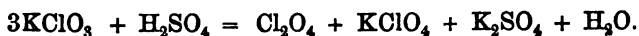
Analytical Reactions (Tests).

First Analytical Reaction.—To solution of a chlorate (e.g. chlorate of potassium) add solution of nitrate of silver; no precipitate falls, showing that the chlorine must be per-

forming different functions from those it possesses in chlorides. Evaporate the solution to dryness, and place the residue in a small dry test-tube, or at once drop a fragment of a chlorate into a test-tube, and heat strongly; oxygen is evolved, and may be recognised by its power of rekindling an incandescent match inserted in the tube. Boil the residue with water, and again add solution of nitrate of silver; a white precipitate falls, having all the characters of chloride of silver, as described under hydrochloric acid.

This is a trustworthy test, and, even omitting the recognition of the oxygen, may be applied in the detection of small quantities of chlorates.

Second Analytical Reaction.—To a fragment of a chlorate add two or three drops of strong sulphuric acid; an explosive gas (Cl_2O_4) is evolved, somewhat resembling chlorine in odour, but possessing a deeper colour than that element.



Warm the upper part of the test-tube to 150° or 200°F. , or introduce a hot wire; a sharp explosion ensues, due to decomposition of the gas, peroxide of chlorine, into its elements.

Third Analytical Reaction.—Heat a small fragment of a chlorate with hydrochloric acid; a yellowish-green explosive gas, termed *euchlorine*, is evolved. Its colour is deeper than that of chlorine, hence the name (from *eu*, well, and *χλωρὸς*, *chlōros*, green). In odour it resembles chlorine, and is probably a mixture of that element with one of the oxides of chlorine.

Fourth Analytical Reaction.—Direct the blowpipe-flame on to charcoal until a spot is red-hot, and then place on the spot a fragment of a chlorate; deflagration ensues as with nitrates.

Bromates.

Bromates are salts closely resembling chlorates and iodates. The formula of Bromic Acid is HBrO_3 .

Iodates.

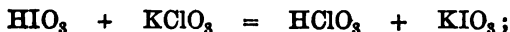
Iodic Acid (HIO_3).—Iodine is boiled in a flask with five times its weight of the strongest nitric acid (sp. gr. 1.5), in a fume-cupboard, until all action ceases. On cooling, iodic acid separates in small pyramidal crystals. These are removed, the residual liquid evaporated to dryness to remove excess of nitric acid, the residue and the first crop dissolved in a small quantity of boiling water and the solution set aside to crystallize.

Iodate of Potassium (KIO_3).—Powder together equal weights of iodine and chlorate of potassium; to the mixture add twice its weight of water and about one-eighth of its weight of nitric acid; warm the whole until iodine disappears and evaporate quite to dryness over a water-bath. The residue dissolved in water forms "Solution of Iodate of Potash," B. P. It contains a little nitrate of potassium.

In this reaction the small quantity of nitric acid furnishes corresponding amounts of nitrate of potassium and chloric acid. The chloric acid with iodine gives iodic acid and chlorine; thus:—

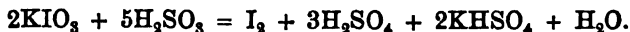


The iodic acid and some chlorate of potassium then yield chloric acid and iodate of potassium,



and the two reactions alternate until the whole of the iodine has displaced the whole of the chlorine.

Iodate of potassium and sulphurous acid decompose each other with elimination of iodine (or with formation of a blue colour, if starch be present). Sulphurous acid occurring as an impurity in acetic and other acids may thus be detected.



Ferric Iodate, or rather *Oxyiodate* ($\text{Fe}_2\text{O}_4\text{IO}_3, 8\text{H}_2\text{O}$), is precipitated on adding solution of ferric chloride to solution of iodate of potassium.

QUESTIONS AND EXERCISES.

460. How may hypochlorous acid be formed ?
461. What are the relations of hypochlorous acid to common bleaching-powder ?
462. By what reaction is chlorine eliminated from hypochlorites ?
463. State the general reaction by which chlorates are formed.
464. Give details of the preparation of chlorate of potassium.
465. Mention the properties of chlorate of potassium.
466. What decompositions occur when chlorate of potassium is heated ?
467. Find the molecular weight of chlorate of potassium.
468. What weight of oxygen is yielded when 1 oz. of chlorate of potassium is completely decomposed ? and how much chloride of potassium remains ?
469. One hundred cubic inches of oxygen, at 60° F. and barometer at 30 inches, weighing 34.203 grains, and 1 gallon containing 277½ cubic inches, what weight of chlorate of potassium will be required to yield 10 gallons of the gas ?—*Ans.* 5½ oz.
470. How many cubic inches of oxygen are producible from 1 oz. of chlorate of potassium ?
471. Calculate the weight of chlorate of potassium theoretically obtainable from 100 parts of chloride.
472. How is perchloric acid prepared ?
473. Enumerate the chlorine acids.
474. How may the presence of chlorides in chlorates be demonstrated ?
475. Mention the tests for chlorates.
476. Give the formula of peroxide of chlorine.
477. What is euchlorine ?
478. How may iodic acid be made ?
479. Describe the preparation of iodate of potassium.

ACETIC ACID AND OTHER ACETATES.

Formula of Acetic Acid $\text{HC}_2\text{H}_3\text{O}_2$, or HA . Molecular weight 60.

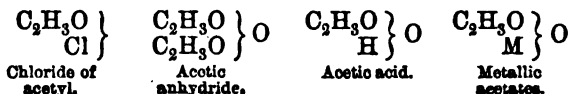
Source.—Acetic acid is said to occur naturally in certain plant-juices and animal fluids in minute proportions, but otherwise is an artificial product. Much is furnished by the destructive distillation of wood ; hence the term *pyroligneous acid* for the crude product, a hybrid word from *πῦρ*, *pûr*, fire,

and *lignum*, wood. This impure product neutralized by carbonate of sodium, the whole evaporated and the residue gently heated to drive off volatile tarry matters, gives acetate of sodium, which after recrystallization furnishes by distillation with oil of vitriol and water acetic acid in a fair state of purity. In Germany and France large quantities of acetic acid are made by the spontaneous oxidation of the alcohol in inferior wines, in the presence, according to Pasteur, of a plant-ferment termed *mycoderma aceti*; hence the *white-* and *red-wine vinegar* (*vinegar*, from the French *vin*, wine, and *aigre*, sour). In England also the domestic form of acetic acid (brown vinegar) has a similar origin: infusion of malt and unmalted grain is fermented; and the resulting oxidation of its sugar, instead of being arrested when the product is an alcoholic liquid, a sort of beer, is allowed to go on to the next stage, acetic acid; it usually contains from 3 to 6 per cent. of real acetic acid ($\text{HC}_2\text{H}_3\text{O}_2$).

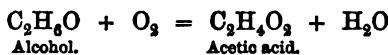
Vinegars.—The official vinegar (*Acetum*, B. P.) contains $5\frac{1}{2}$ (5.4) per cent. of acid. The so-called Vinegar of Cantharides (*Acetum Cantharidis*, B. P.) is a solution of the active principle of cantharides in very strong acetic acid, not in vinegar. The Vinegar of Squill (*Acetum Scillæ*, B. P.) is also a solution of the active principle of squill in dilute acetic acid, not in true vinegar. In the Pharmacopœia, vinegar, except its use for its own sake, is only employed in the preparation of *Emplastrum Cerati Saponis*.

The Acetic Radical.—The group of elements represented by the formula $\text{C}_2\text{H}_3\text{O}_2$ is that characteristic of acetic acid and other acetates, and may, for convenience of study, be assumed to be an acidulous univalent radical. It has not been isolated, unless, indeed, a compound of similar composition, resulting from the action of peroxide of barium on acetic anhydride, is the radical in question.

Acetyl.—The characteristic grouping in acetates ($\text{C}_2\text{H}_3\text{O}_2$) is frequently considered to contain, rather than to be, a radical—the radical $\text{C}_2\text{H}_3\text{O}$, termed *acetyl*. Acetates yield a body having the composition $\text{C}_2\text{H}_3\text{OCl}$, which is regarded as chloride of acetyl; from this may be obtained acetic anhydride ($\text{C}_4\text{H}_6\text{O}_3$), which by absorbing water becomes acetic acid.



The relation of acetic acid to alcohol will be evident from the following equation representing, empirically, the formation of the acid :—



Acetates in aqueous solution are liable to decomposition. In solution of acetate of morphia a myceloid growth occasionally forms, acetic acid disappears, and morphia is deposited. Solution of acetate of ammonium is liable to a similar change, gradually becoming alkaline.

Synthetical Reaction.

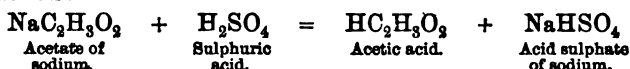
Acetic Acid.

To a few grains of acetate of sodium in a test-tube add a little water and some sulphuric acid, and heat the mixture; acetic acid is evolved, and may be condensed by a bent tube adapted to the test-tube by a cork in the usual way.

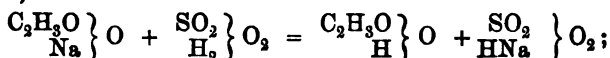
Acetic Acid.—This is the process by which acetate of sodium or calcium (the neutralized products of the distillation of wood) is made to yield acetic acid on the large scale. As with nitric and hydrochloric acids, the loose term “acetic acid” is that usually applied to aqueous solutions of acetic acid. The *Acidum Aceticum*, B. P., contains nearly 33 per cent. of real acid, that is, of $\text{HC}_2\text{H}_3\text{O}_2$; for it contains only 28 per cent. of *acetic anhydride* ($\text{C}_4\text{H}_6\text{O}_3$)—still occasionally, though somewhat obscurely, termed *anhydrous acetic acid*. *Acidum Aceticum Dilutum*, B. P., contains about $4\frac{1}{4}$ per cent. of $\text{HC}_2\text{H}_3\text{O}_2$. Glacial acetic acid ($\text{HC}_2\text{H}_3\text{O}_2$) contains no water. It solidifies to a crystalline mass at temperatures below 63°F ., hence the appellation *glacial* (from *glacies*, ice). Good commercial glacial acetic acid (*Acidum Aceticum Glaciale*, B. P.) does not contain more than 1 per cent. of water, corresponding to 84.15 per cent. of acetic anhydride; it solidifies at 34°F ., and again liquefies at 48° : its specific gravity is 1.065. Although water is lighter than this acetic acid, yet the addition of water at first renders the acid heavier; evidently therefore condensation, or contraction in bulk, occurs on mixing the liquids: after 10 per cent. has been added, the addition of more water produces the usual effect of dilution of a heavy liquid by a lighter—namely, reduction of relative

weight. This matter will be better understood after the subject of specific gravity has been studied. Glacial acetic acid mixes readily with most oils.

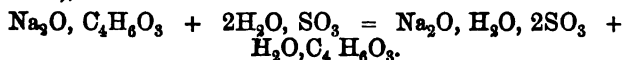
The following equation is expressive of the foregoing reaction:—



or, assuming the existence of acetyl ($\text{C}_2\text{H}_3\text{O}$) in acetic acid, and a corresponding radical sulphuryl (SO_2) in sulphuric acid,



or, thirdly, on the assumption that salts contain the oxide of a basylous radical united with the anhydride of an acid (the old view under which such names as acetate of soda were formed),



Note on the Constitution of Salts.

Which of these three equations, or, more broadly, which of the three views of the constitution of salts illustrated by the equations, is correct, it is impossible to say. Whether it is $\text{C}_2\text{H}_3\text{O}_2$, $\text{C}_2\text{H}_3\text{O}$, or $\text{C}_4\text{H}_6\text{O}_3$ which migrates from one acetic compound to another, whether it is SO_4 , SO_2 , or SO_3 which migrates from one sulphuric compound to another, and so on with other acidulous groupings, cannot at present be determined. There are strong objections to each view; and possibly neither is right. Either the given radicals cannot be isolated; or application of the forces of heat, light, and electricity do not confirm views arrived at by the results of operations with the chemical force; or a salt comes to be regarded as having so large a number of constituent parts that the view, however true, breaks down in practice from the sheer inability of the mind to grasp the complicated analogies involved. Yet for the purposes of description, study, and conversation some system must be adopted. Let the first, then, be generally taken, over-reliance on it being checked by the use of general instead of special names for the hypothetical radicals, and other systems be employed in certain cases. (See also p. 331.)

Impurity.—Acetic acid sometimes contains sulphurous acid. The methods by which this impurity is detected are described on pages 344 and 357.

Analytical Reactions (Tests).

First Analytical Reaction.—To an acetate add sulphuric acid and heat the mixture; acetic acid, recognised by its odour, is evolved.

Note 1.—Iodine, sulphurous acid, and other substances of powerful odour mask that of acetic acid; they must be removed, therefore, usually by precipitation or oxidation, before applying this test.

Note 2.—It will be noticed that this reaction is identical with the previous one; it has synthetical or analytical interest, according to the object and method of its performance.

Second Analytical Reaction.—Repeat the above action, a few drops of spirit of wine being first added to the acetate; acetic ether (acetate of ethyl, $C_2H_5C_2H_5O_2$), also of characteristic odour, is evolved.

The basylous radical ethyl (C_2H_5) will be referred to subsequently.

Third Analytical Reaction.—Heat a fragment of a dry acetate in a test-tube, and again notice the odour of the gaseous products of the decomposition; among them is *acetone* (C_3H_6O), the smell of which is characteristic. Carbonate of the metal remains in the test-tube.

Fourth Analytical Reaction.—To a solution of an acetate, made neutral by the addition of acid or alkali, as the case may be, add a few drops of neutral solution of perchloride of iron; a deep-red liquid results, owing to the formation of ferric acetate ($Fe_2(C_2H_3O_2)_6$).

Analytical Note.—It will be noticed that the formation of characteristic precipitates, the usual method of removing radicals from solution for recognition, is not carried out in the qualitative analysis of acetates. This is because all acetates are soluble. Acetate of silver ($AgC_2H_3O_2$) and mercurous acetate ($Hg_2C_2H_3O_2$) are only sparingly soluble in cold water,

but the fact can seldom be utilized in analysis. Hence peculiarities of colour and odour, the next best characters on which to rely, are adopted as means by which acetates may be detected. Acetates, like other organic compounds, char when heated to a high temperature.

Note on Anhydrides.—Up to this point the student has regarded an anhydride as a body derived from an acid by removal of the *whole* of the hydrogen of the acid, together with as much of its oxygen as with the hydrogen forms water. This definition will scarcely apply to acetic anhydride, and must therefore be somewhat qualified. An *anhydride* is derived from an acid, the acid having lost the whole of its *basylous* hydrogen and so much oxygen as is necessary to form water with that hydrogen. Anhydrides are obtained by heating acids, and by other methods.

QUESTIONS AND EXERCISES.

480. What is the formula of acetic acid ?
481. State the relation of acetic acid to other acetates.
482. What is the molecular weight of acetic acid ?
483. Name the sources of acetic acid.
484. What is pyroligneous acid ?
485. From what compound is the acetic acid of foreign and English vinegar immediately derived ?
486. How much real acid is contained in official vinegar ?
487. What is the nature of the "Vinegars" of Pharmacy ?
488. How may acetic acid be obtained from acetate of sodium ?
489. How much real acid is contained in the official acetic acid ?
490. Mention the strength of commercial glacial acetic acid.
491. Give three or more views of the constitution of acetates, illustrating each by formulæ.
492. Enumerate the tests for acetates.

HYDROSULPHURIC ACID AND OTHER SULPHIDES.

Formula of Hydrosulphuric Acid H_2S . Molecular weight 34.

Source and Varieties of Sulphur.—The acidulous radical of *hydrosulphuric acid*, *sulphydric acid*, or *sulphuretted hydrogen* and other sulphides, is the element sulphur (S). It occurs in

nature in combination with metals, as already stated in describing the ores of some of the metals, and also in the free state. Most of the sulphur used in medicine is imported from Sicily, where it occurs chiefly associated with blue clay. It is purified by fusion, sublimation, or distillation. Melted and poured into moulds, it constitutes a crystalline mass termed *roll sulphur*. If distilled and the vapour carried into large chambers, so that it may be rapidly condensed, the crystals are so minute as to give the sulphur a pulverulent character; this is *sublimed sulphur* (*Sulphur Sublimatum*, B. P.), or *flowers of sulphur*: the same washed constitutes *Sulphur Lotum*, U. S. P. The third common form, *milk of sulphur*, will be noticed subsequently. Sulphur also occurs in nature in combination as a constituent of animal and vegetable tissues, as sulphurous acid gas (SO_2) in volcanic vapours, and as sulphuretted hydrogen in some waters, as those of Harrogate. *Plastic sulphur* is one of the allotropic varieties of the element, obtained on heating sulphur considerably beyond its melting-point, and pouring into cold water.

Quantivalence.—Sulphur is sexivalent, as seen in sulphuric anhydride (SO_3) a substance which will be noticed under sulphuric acid. It also occasionally exhibits quadrivalent (SO_2) and still oftener bivalent affinities (H_2S).

Molecular Weight.—At very high temperatures sulphur follows the rule that, under similar conditions of heat and pressure, atomic weights (in grammes, grains, etc.) of volatile elements occupy equal volumes of vapour; its formula therefore is S_2 , and molecular weight 64. At lower temperatures the volume weighs three times as much as it should do if following usual laws, and then the molecule would appear to contain six atoms (S_6).

Acid Salts.—Sulphur (S'') being the first acidulous radical of bivalent activity met with in these sections on acids, it is desirable here to draw attention to a new class of salts to which such a radical will generally give rise. These are acid salts, which are intermediate between normal salts and acids. Univalent radicals with an atom of hydrogen give an acid, and with an atom of other basylous radicals an ordinary or normal salt. But bivalent radicals, from the fact that they give with two atoms of hydrogen an acid, and with two atoms of univalent metals a normal salt, may obviously give intermediate bodies containing one atom of hydrogen and one atom of metal; these are appropriately termed acid salts: they

are neither normal acids nor normal salts, but *acid salts*. (Examples:— KHCO_3 , NaHSO_4 , $\text{KHC}_4\text{H}_4\text{O}_6$, Na_2HPO_4 , CuHAsO_3 , CaH_2PO_4 .) Whether or not these salts give an acid reaction with blue litmus paper depends on the strength of the respective radicals. Usually they do redden the test-paper, but sometimes not; thus the acid sulphide or sulphhydrate of potassium (KHS), of sodium (NaHS), or ammonium (AmHS) has alkaline properties.*

The chemical analogy between sulphur and oxygen, already once alluded to (p. 196), is further illustrated by the compounds just mentioned. Sulphur is also closely related to the rarer element selenium. Thus we have SeO_2 as well as SO_2 , H_2SeO_3 (selenious acid) as well as H_2SO_3 (sulphurous acid), H_2SeO_4 (selenic acid) as well as H_2SO_4 (sulphuric acid). The rare element tellurium also seems to have similar analogies. The four hydrogen compounds of the group have the formulæ H_2O , H_2S , H_2Se , H_2Te .

Synthetical Reactions.

Sulphuretted Hydrogen.

First Synthetical Reaction.—The preparation of sulphuretted hydrogen.—This operation was described on page 102, and probably has already been studied by the reader.

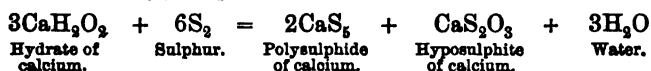
Precipitated Sulphur.

Second Synthetical Reaction.—Prepare the variety of the radical of sulphides known as *Precipitated Sulphur* (*Sulphur Præcipitatum*, B. P.), or *Milk of Sulphur*, by boiling a few grains of flowers of sulphur (5 parts) with slaked lime (3 parts) and some water (20 parts) in a test-tube (larger quantities in an evaporating-basin), filtering, and (reserving a small portion of the filtrate) adding dilute hydrochloric acid until the well-stirred milk-like liquid has a faint acid reaction on test-paper; sulphur is precipitated, and may be collected

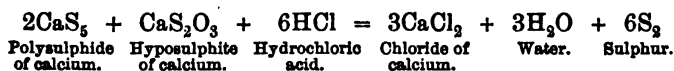
* Some chemists regard these sulphhydrates as compounds of basylous radicals with HS, a univalent grouping termed hydrosulphyl (persulphide of hydrogen, H_2S_2), just as hydrates are similarly viewed as compounds of the univalent radical hydroxyl (HO) (peroxide of hydrogen, H_2O_2).— H_2S becoming HHS or HHs (hydrosulphylide of hydrogen), and H_2O becoming HHO or HHo (hydroxylide of hydrogen).

on a filter, washed, and dried (at about 120° F.). Excess of acid must be avoided, or some hydrosulphyl, the liquid polysulphide of hydrogen (H_2S_2), will be formed, probably causing the particles of sulphur to aggregate to a gummy mass.

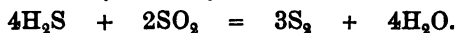
This is the process of the Pharmacopœia. Polysulphide of calcium and hyposulphite of calcium are formed :—



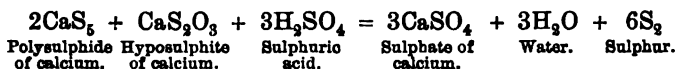
On adding the acid, both salts are decomposed and, after an intermediate reaction, sulphur separates :—



The polysulphide of calcium yields sulphuretted hydrogen and milk-white sulphur on the addition of acid. The hyposulphite of calcium then yields sulphurous acid gas as well as yellowish sulphur. The gases react and give sulphur and water, very little sulphuretted hydrogen escaping. A little pentathionic acid (*see* Index) is also said to be formed.



Calcareous Precipitated Sulphur. The old "Milk of Sulphur."—To a sulphur solution prepared as before (or to the reserved portion) add a little dilute sulphuric acid; the precipitate is in this case largely mixed with sulphate of calcium :—



Place a little of each of these specimens of precipitated sulphur with a drop of the supernatant liquid on a strip of glass, cover each spot with a piece of thin glass, and examine the precipitates under a microscope; the pure sulphur will be found to consist of minute grains or globules, the calcareous to contain comparatively large crystals (sulphate of calcium).

Notes.—A large proportion of the precipitated sulphur met with in trade is, in England, still thus mixed with sulphate of

calcium, most specimens containing two-thirds of their weight of that substance. Many purchasers, indeed, are so accustomed to the satiny appearance of the mixed article as to regard real sulphur with suspicion, sometimes refusing to purchase it. The mixed article is, certainly, more easily miscible with aqueous liquids: but it has long been discarded from the Pharmacopœias of Great Britain, and, on the whole, is scarcely worthy of a prominent place in the pharmacy of 1880.

Many English pharmacists have ceased to sell precipitated sulphur which yields a white ash (the anhydrous sulphate) when a little is burnt off on the end of a table-knife or spatula. (No more damage is done to the steel than a rub on a knife-board will remove.)

To ascertain, exactly, the amount of sulphate of calcium in a specimen of calcareous precipitated sulphur, place a weighed quantity in a tared crucible and heat till no more vapours are evolved. The weight of the residual anhydrous sulphate of calcium ($\text{CaSO}_4 = 136$), with one-fourth thereof added, is the amount of crystalline sulphate of calcium ($\text{CaSO}_4, 2\text{H}_2\text{O} = 172$) present in the original quantity of calcareous sulphur.

Analytical Reactions (Tests).

To a sulphide add a few drops of hydrochloric acid; sulphuretted hydrogen will probably be evolved, well known by its smell. If the sulphide is not acted upon by the acid, or if free sulphur be under examination, mix a minute portion with a fragment of solid caustic potash or soda, and fuse on a silver coin or spoon. When cold, place a drop of dilute hydrochloric acid on the spot, sulphuretted hydrogen is evolved, and a black stain, due to sulphide of silver (Ag_2S), left on the coin.

Other sulphur reactions may be adopted as tests; but the above are sufficient for all ordinary purposes. The most convenient reagent for detecting a sulphide in solution of ammonia is ammonio-sulphate of copper, which gives a black precipitate of sulphide of copper if a sulphide be present.

The *Iodide of Sulphur* (S_2I_2) has been mentioned under "Iodine." A *chloride* (S_2Cl_2) and *bromide* (S_2Br_2) may also be formed from their elements. A mixture of sulphur and

chloride of sulphur is sometimes met with under the name of *hypochloride of sulphur*.

QUESTIONS AND EXERCISES.

493. In what forms does sulphur occur in nature?
 494. State the modes of preparation of the three chief commercial varieties of sulphur.
 495. To what extent does the atom of sulphur vary in valence?
 496. State the relations of acid salts to acids and to normal salts.
 497. Define sulphides and sulphydrates.
 498. Describe the preparation of sulphuretted hydrogen.
 499. What are the characters of pure precipitated sulphur?
 500. Give equations explanatory of the reactions which occur in precipitating sulphur according to the official process.
 501. Describe the microscopic test for calcareous precipitated sulphur.
 502. Mention a ready physical method of detecting sulphate of calcium in precipitated sulphur.
 503. Mention the tests for sulphides, and the character by which sulphuretted hydrogen is distinguished from other sulphides.
 504. How are sulphides insoluble in acids tested for sulphur?
 505. Give a method for the detection of a trace of sulphur in solution of ammonia.
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SULPHUROUS ACID AND OTHER SULPHITES.

Formula of sulphurous acid H_2SO_3 . Formula of sulphurous acid gas or sulphurous anhydride, commonly termed sulphurous acid, SO_2 . Molecular weight of sulphurous acid 82; of the anhydride, 64.

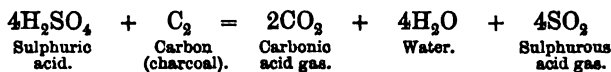
When sulphur is burnt in the air it combines with oxygen and forms sulphurous acid gas (SO_2), more correctly termed sulphurous anhydride, or commonly, but erroneously, sulphurous acid. It is a pungent, colourless gas, readily liquefied on being passed through a tube externally cooled by a *freezing-*

mixture composed of two parts of well-powdered ice (or, better, snow) with one part of common salt. If sulphurous acid gas becomes moist or is passed into water, heat is evolved and true sulphurous acid (H_2SO_3) formed. The latter body may be obtained in crystals; but it is very unstable, and hence the properties of the sulphurous radical must be studied under the form of some other sulphite; sulphite of calcium (CaSO_3), or sulphite of sodium (Na_2SO_3), may be used for the purpose.

Quantivalence.—The radical of the sulphites is bivalent (SO_3''), and hence forms acid sulphites, such as acid sulphite of potassium (KHSO_3), and normal sulphites, such as sulphite of sodium (Na_2SO_3).

Note on Nomenclature.—The sulphites are so named from the usual rule, that salts corresponding with acids whose names end in *ous* have a name ending in *ite*. They are generally made by passing sulphurous acid gas over moist oxides or carbonates; in the latter case carbonic acid gas escapes.

Synthetical Reaction.—To a few drops of sulphuric acid in a test-tube add a piece of charcoal and apply heat; sulphurous acid gas is evolved, and may be conveyed by a bent tube into a small quantity of cold water in another test-tube. Larger quantities may be made in a Florence flask. The product is the *Acidum Sulphurosum*, B. P. It is said to contain, if saturated, nearly 12 (11·79) per cent. of sulphurous acid (H_2SO_3), or about 9 (9·2) per cent. of the gas (SO_2). The process is also that described in the *Pharmacopoeia*, except that the gas is purified by passing through a small wash-bottle before the final collection.



Sulphurous acid gas may also be made by boiling copper, mercury, or iron with sulphuric acid, sulphate of the metal being formed. Also by boiling sulphur with sulphuric acid.



If in this process the water were replaced by solutions of, or solid, metallic oxides or carbonates, sulphites of the various metals would be formed. The formula of Sulphite of Sodium (*Sodii Sulphis*, U. S. P.) is $\text{Na}_2\text{SO}_3 \cdot 7\text{H}_2\text{O}$; of the Bisulphite NaHSO_3 . Under the name of *antichlor* the former is used for removing traces of chlorine from paper pulp. The so-called *Bisulphite of Lime*, used by brewers for retarding or arresting fermentation and oxidation, and for various antiseptic purposes, is a solution of sulphite of calcium (CaSO_3) in free sulphurous acid (H_2SO_3), and is made by passing sulphurous acid gas (SO_2) into thin milk of lime. Its specific gravity varies from 1.050 to 1.070, and its potential strength of anhydride (SO_2) from 4 to 6 per cent.

Analytical Reactions (Tests).

First Analytical Reaction.—To a sulphite (of sodium, for instance,—made by passing sulphurous acid into solution of carbonate of sodium) add a drop or two of dilute hydrochloric acid; sulphurous acid gas escapes, recognised by a peculiarly pungent smell.

This smell is the same as that evolved on burning lucifer matches that have been tipped with sulphur. It is due, probably not to the gas (SO_2), but to sulphurous acid (H_2SO_3) formed by the union of sulphurous acid gas with either the moisture of the air or that on the surface of the mucous membrane of the nose. The gas is highly suffocating.

Second Analytical Reaction.—To a sulphite add a little water, a fragment or two of zinc, and then hydrochloric acid; sulphuretted hydrogen will be evolved, known by its putrid odour and action on a piece of paper placed like a cap on the mouth of the test-tube and moistened with a drop of solution of acetate of lead, black sulphide of lead being formed. Sulphurous acid may be detected in acetic acid, or in hydrochloric acid, by this test.



Other Analytical Reactions.

To solutions of neutral sulphites add nitrate or chloride of barium, chloride of calcium, or nitrate of silver; in each case

white sulphites of the various metals are precipitated. The barium sulphite is soluble in weak hydrochloric acid; but if a drop or two of chlorine-water is first added, barium sulphate is formed, which is insoluble in acids. The other precipitates are also soluble in acids. The silver sulphite is decomposed on boiling, sulphuric acid being formed, and metallic silver set free.

To recognise the three radicals in an aqueous solution of sulphides, sulphites, and sulphates, add chloride of barium, filter, and wash the precipitate. In the filtrate sulphides are detected by the sulphuretted hydrogen evolved on adding an acid. In the precipitate, sulphites are detected by the odour of sulphurous acid produced on adding hydrochloric acid, and sulphates by their insolubility in the acid.

QUESTIONS AND EXERCISES.

506. What are the differences between sulphurous acid and sulphurous acid gas, sulphites and acid sulphites?

507. State the characters of sulphurous acid gas.

508. How is the official Sulphurous Acid prepared?

509. By what tests may sulphurous acid be recognised in acetic acid?

510. Give a method by which sulphites may be detected in presence of sulphides and sulphates.

SULPHURIC ACID AND OTHER SULPHATES.

Formula of Sulphuric Acid H_2SO_4 . Molecular weight 98.

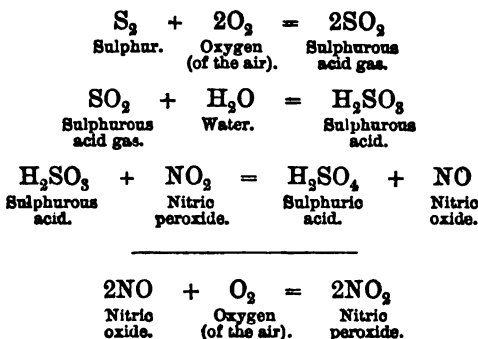
Many sulphates occur in nature; but the common and highly important hydrogen sulphate, sulphuric acid, is made artificially.

Preparation of the acid. General nature of the process.—Sulphur itself, or sometimes the sulphur in iron pyrites, is first converted into sulphurous acid gas by burning in air, and this gas, by moisture and oxygen, into sulphuric acid ($\text{SO}_2 + \text{H}_2\text{O} + \text{O} = \text{H}_2\text{SO}_4$).

Details of the process.—The oxygen necessary to oxidize the sulphurous acid gas cannot directly be obtained from air, but

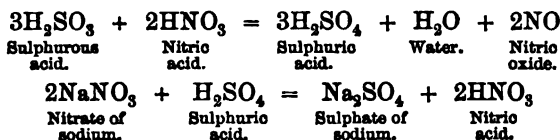
indirectly, the agency of nitric oxide (NO) being employed—this gas becoming nitric peroxide (NO₂) by action of the air, and the nitric peroxide again becoming nitric oxide by the action of the sulphurous acid gas, and so on. A small quantity of nitric oxide gas will in this way act as carrier of oxygen from the air to very large quantities of sulphurous acid.

The following equations represent the successive steps:—



On the large scale the sulphurous acid gas is produced by burning sulphur in furnaces; it is carried, together with the nitric vapours, by flues into leaden chambers, where jets of steam supply the necessary moisture; the steam also, condensing, prevents other reactions. The resulting dilute sulphuric acid is concentrated by evaporation in leaden, and finally in glass or platinum vessels.

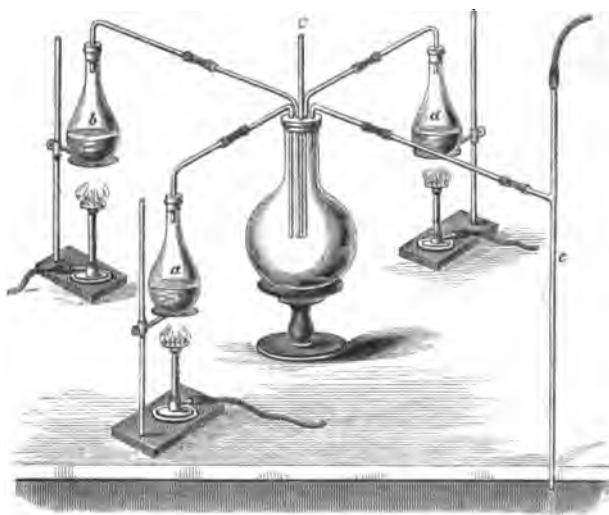
The nitric oxide is in the first instance obtained from nitric acid, and this from nitrate of potassium or of sodium by the action of a small quantity of the sulphuric acid of a previous operation.



Other processes.—Sulphuric acid may be obtained by other processes, as by distilling the sulphate of iron resulting from the natural oxidation of iron pyrites by air; but it is seldom

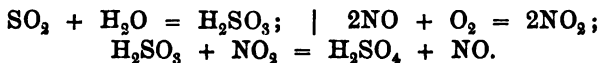
so made at the present day. The sulphate of iron was formerly called *green vitriol* (p. 157), and the distilled product *oil of vitriol*; the latter in allusion to its consistence and origin.

Fig. 39.



EXPERIMENTAL MANUFACTURE OF SULPHURIC ACID.

Experiment.—For purposes of practical study a small quantity may be made, as shown in fig. 39, by passing, *a*, sulphurous acid gas (p. 356), *b*, nitric oxide in small quantity (p. 335), *c*, air (forced through by aid of bellows or a gas-holder, or drawn through by a *water-aspirator*, *e*), and, occasionally, *d*, steam (generated in a Florence flask) through glass tubes, nearly to the bottom of a two- or three-quart flask.



A slow current of sulphurous acid gas, air, and steam, and a small quantity of nitric oxide, will furnish, in the course of a few minutes, enough sulphuric acid for recognition by the first of the following analytical reactions.

Purification.—Sulphuric acid may contain arsenic, nitrous compounds, and salts. Arsenic may be detected by the hydrogen test (p. 192), nitrous compounds by powdered sulphate of iron (which acquires a violet tint if they are present), and salts by the residue left on boiling a little to dryness in a crucible in a fume-chamber. If only nitrous compounds are present, the acid may be purified by heating with about half per cent. of sulphate of ammonium—water and nitrogen being produced (Pelouze). If arsenic occurs, boil with a small quantity of hydrochloric acid, which converts the arsenic into volatile chloride of arsenicum; or heat with a little nitric acid, which converts arsenious (As_2O_3) into arsenic anhydride (As_2O_5), then add sulphate of ammonium, and distil in a retort containing pieces of quartz and heated by an annular-shaped burner (to prevent "bumping"—see page 324). The arsenic anhydride remains in the retort. (Arsenious anhydride would be carried over with the sulphuric-acid vapours.) By distillation the acid is purified from salts (such as NaHSO_4) which are not volatile.

Quantivalence.—The sulphuric radical being bivalent (SO_4''), acid as well as normal sulphates may exist. Acid sulphate of potassium (KHSO_4) is an illustration of the former, sulphate of sodium (Na_2SO_4) of the latter; *double sulphates* may also occur, such as that of potassium and magnesium ($\text{K}_2\text{SO}_4, \text{MgSO}_4, 6\text{H}_2\text{O}$). Sulphates generally contain water of crystallization.

Pure sulphuric acid (H_2SO_4) is of specific gravity 1.848. The best "oil of vitriol" of commerce, a colourless liquid of oily consistence, is of specific gravity 1.843, and contains 96.8 per cent. of real acid (H_2SO_4). This is the *Acidum Sulphuricum*, B. P. The *Acidum Sulphuricum Dilutum*, B. P., sp. gr. 1.094, contains about $13\frac{1}{2}$ (13.64) per cent. of acid (H_2SO_4); and the *Acidum Sulphuricum Aromaticum*, B. P., a dilute acid in which is dissolved the soluble aromatic parts of cinnamon and ginger, also contains nearly $13\frac{1}{2}$ (13.36) per cent. of acid (H_2SO_4). It has been stated that in this preparation the acid and alcohol form some sulphovinic acid, but the author has been unable to detect the latter. There are some definite compounds of sulphuric acid with water; the first ($\text{H}_2\text{SO}_4, \text{H}_2\text{O}$) may be obtained in crystals.

Sulphuric anhydride (SO_3) occurs in white crystals having no acid properties. It is made by distilling sulphuric acid with phosphoric anhydride ($3\text{H}_2\text{SO}_4 + \text{P}_2\text{O}_5 = 2\text{H}_3\text{PO}_4 + 3\text{SO}_3$).

It appears to unite with sulphuric acid and some other normal sulphates to form compounds (R'_2SO_4 , SO_3) resembling in constitution red chromate of potassium. The fuming sulphuric acid ($H_2SO_4 \cdot SO_3$) formerly made at Nordhausen in Saxony seems to be such a body.

Note.—Sulphuric acid is a most valuable compound to all chemists and manufacturers of chemical substances. By its agency, direct or indirect, many, if not most, chemical transformations are effected. To describe all its uses would be to write a work on chemistry.

Analytical Reactions (Tests).

First Analytical Reaction.—To solution of a sulphate add solution of a barium salt; a white precipitate of sulphate of barium ($BaSO_4$) falls. Add nitric acid and boil the mixture, the precipitate does not dissolve.

This reaction is as highly characteristic of sulphates as it has been stated to be of barium salts (*vide* page 110). The only error likely to be made in its application is that of overlooking the fact that nitrate and chloride of barium are less soluble in strong acid than in water. On adding the barium salt to the acid liquid, therefore, a white precipitate may be obtained, which is simply the nitrate or chloride of barium. The appearance of such a precipitate differs considerably from that of the barium sulphate; hence a careful operator will not be misled. Should any doubt remain, water should be added, which will dissolve the nitrate or chloride, but not affect the sulphate.

Second Analytical Reaction.—Mix a fragment of an insoluble sulphate ($BaSO_4$ *e.g.*) with carbonate of potassium or of sodium, or, better, with both carbonates, and fuse the mixture in a small crucible. Digest the residue when cold, in water, and filter; the filtrate may be tested for the sulphuric radical.

This is a convenient method of qualitatively analysing insoluble sulphates, such as those of barium and lead.

Third Analytical Reaction.—Mix a fragment of an insoluble sulphate with a little alkaline carbonate on a piece of charcoal, taking care that some of the charcoal-dust is included in the mixture. Heat the little heap in the blowpipe-flame until it

fuses, and, when cold, add a drop of acid; sulphuretted hydrogen is evolved, recognised by its odour.

This is another process for the recognition of insoluble sulphates. Other preparations of sulphur, and sulphur itself, give a similar result. It is therefore rather a test for sulphur and its compounds than sulphates only; but the absence of other salts can generally, if necessary, be previously determined.

Note.—The presence of the sulphuric radical in a solution having been proved by the above reactions, its occurrence as the normal sulphate of a metal is demonstrated by the neutral, or nearly neutral, deportment of the liquid with test paper, and the detection of the metal—its occurrence as sulphuric acid or an acid sulphate by the sourness of the liquid to the taste, and the effervescence produced on the addition of a carbonate.

Antidote.—In cases of poisoning by strong sulphuric acid solution of carbonate of sodium (common washing-soda), magnesia and water, etc., may be administered as antidotes.

QUESTIONS AND EXERCISES.

511. What is the formula of sulphuric acid, and what its molecular weight?

512. How is it related to other sulphates?

513. Write a short article on the manufacture of sulphuric acid, giving either diagrams or equations.

514. How may nitrous compounds be detected in, and eliminated from, sulphuric acid?

515. State the method by which the presence of arsenic is detected in sulphuric acid, and explain the process by which it may be removed.

516. Define sulphates, acid sulphates, and double sulphates.

517. What percentage of real acid is contained in commercial oil of vitriol?

518. State the strength of the official "diluted" and "aromatic" sulphuric acid.

519. By what process is sulphuric anhydride obtained from Nordhausen sulphuric acid?

520. Explain the reactions which occur in testing for sulphates.

521. Ascertain by calculation the weight of oil of vitriol (of 96·8 per cent.) necessary for the production of one ton of dry sulphate of ammonium.—*Ans.* 1718 lbs.

522. Name the antidotes in cases of poisoning by strong sulphuric acid.

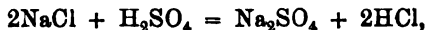
CARBONIC ACID AND OTHER CARBONATES.

Formula of carbonic acid H_2CO_3 . Molecular weight 62.

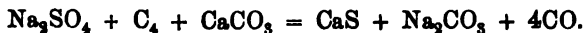
Formula of carbonic acid gas, or carbonic anhydride, commonly termed carbonic acid, CO_2 .

Sources.—Carbonates (compounds containing the grouping CO_3) are very common in nature, the calcium carbonate (CaCO_3) being widely distributed as chalk, limestone, or marble. The hydrogen carbonate, true carbonic acid, is not known, unless, indeed, carbonic acid gas assumes that condition on dissolving in water. Such a solution (see p. 88) changes the colour of blue litmus paper, and the gas does not; this may be because only the true acid (H_2CO_3) affects the litmus, or because the gas (CO_2) cannot come into real contact with the litmus without a medium. From the commonest natural carbonate, carbonate of calcium, is derived the carbonic constituent of the one most frequently used in medicine and the arts generally, carbonate of sodium.

Carbonate of sodium is prepared from the chief natural salt, the chloride. After the chloride has been converted into sulphate (salt-cake) by sulphuric acid (or by sulphurous acid, air, and steam—Hargreave's modification),

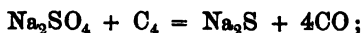


the sulphate is roasted with limestone and small coal, by which carbonate of sodium and sulphide of calcium are formed:—

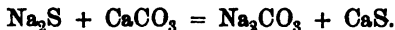


Carbonic oxide gas and a little carbonic acid gas, from the excess of chalk, escape; the residual mass (black ash) is digested in water, in which the carbonate of sodium dissolves, the sulphide of calcium, with a little oxide, remaining insoluble. The solution is evaporated to dryness, and yields crude

carbonate of sodium. This is roasted with a small quantity of sawdust, to convert any caustic soda, resulting from the action of the lime on the carbonate, into normal carbonate. The product is *soda-ash*. Dissolved in water and crystallized, it constitutes the ordinary "soda" used for washing purposes: recrystallized and sometimes ground, it forms the official carbonate of sodium (*Sodæ Carbonas*, B. P.) ($\text{Na}_2\text{CO}_3, 10\text{H}_2\text{O}$). The reaction is rendered more intelligible by regarding it as occurring in two stages:—1st, the reduction of the sulphate of sodium to sulphide by the carbon of the coal,



2nd, the reaction of the sulphide of sodium and carbonate of calcium, giving soluble carbonate of sodium, thus—



The sulphur in the residual sulphide (or, perhaps, oxy-sulphide) of calcium may be recovered by exposure to air, and the subsequent action of hydrochloric acid. Some hypsulphite of calcium (CaS_2O_3) is first formed, and the action of the acid on this and on undecomposed sulphide gives chloride of calcium, water, and sulphur.

Another process.—To a strong solution of common salt, bicarbonate of ammonium is added, when a precipitate of bicarbonate of sodium occurs. The resulting chloride of ammonium may be converted into carbonate by heating with chalk, and the carbonate be more fully carbonated by carbonic acid gas obtained by heating the bicarbonate of sodium, which is thereby reduced to the ordinary neutral carbonate.

Carbonic acid gas (CO_2) is a product of the combustion of all carbonaceous matters. It is constantly exhaled by animals and inhaled by plants, its intermediate storehouse being the atmosphere, throughout which it is equally distributed by *diffusion* (*vide* p. 14) to the extent of about 4 parts in 10,000. A larger proportion than that just mentioned gives to confined air depressing effects, 4 or 5 per cent. rendering the atmosphere poisonous when taken into the blood from the lungs. Carbonic acid, however, may be taken into the stomach with beneficial sedative effects; hence, probably, much of the value of such effervescing liquids as soda-water, lemonade, and solutions of the various granulated preparations and effervescing powders. The gas liquefies on being compressed, and the liquid solidifies on being cooled. Carbonic acid gas is twenty-

two times as heavy as hydrogen, and about half as heavy again as air.

Sulphocarbonates resemble carbonates in constitution, but contain sulphur in place of oxygen.

REACTIONS.

Synthetical and Analytical Reactions.—1. To a fragment of marble in a test-tube add water and then hydrochloric acid; carbonic acid gas (CO_2) is evolved, and may be conveyed into water or solutions of salts by the usual delivery-tube.

This is the process of the British Pharmacopœia, and the one usually adopted for experimental purposes. Passed into carbonate of sodium, the gas gives *Sodæ Bicarbonas*, B. P. (p. 85), and into carbonate of potassium, *Potassæ Bicarbonas*, B. P. (p. 71). On the large scale the gas is prepared from chalk or marble and sulphuric acid, frequent stirring promoting its escape.

2. Pass the gas into lime-water; a white precipitate of carbonate of calcium (CaCO_3) falls. Solution of subacetate of lead may be used instead of, and is perhaps even a more delicate test than, lime-water.

The evolution of a gas on adding an acid to a salt, warming the mixture if necessary, the gas being inodorous and giving a white precipitate with lime-water, is sufficient evidence of the presence of a carbonate. Carbonates in solutions of ammonia, potash, or soda may be detected by the direct addition of solution of lime. Carbonates in presence of sulphites or hyposulphites may be detected by adding acid tartrate of potassium, which decomposes carbonates with effervescence, but does not attack sulphites or hyposulphites.

3. Blow air from the lungs through a glass tube into lime-water; the presence of carbonic acid gas is at once indicated.

The passage of a considerable quantity of normal air through lime-water produces a similar effect. A bottle containing lime-water soon becomes coated with carbonate of calcium from absorption of atmospheric carbonic acid gas.

4. Fill a dry test-tube with the gas, by passing the delivery-tube of the above apparatus to the bottom of the test-tube.

Being rather more than once and a half as heavy as the air (1.529), it will displace the latter. Prove the presence of the gas by pouring it slowly, as if a visible liquid, into another test-tube containing lime-water; the characteristic cloudiness and precipitate are obtained on gently shaking the lime-water.

In testing for carbonates by bringing evolved gas into contact with lime-water, the preparation and adaptation of a delivery-tube may often be avoided by pouring the gas from the generating-tube into that containing the lime-water in the manner just indicated.

5. Pass carbonic acid gas through lime-water until the precipitate at first formed is dissolved. The resulting liquid is a solution of carbonate of calcium in carbonic acid water. Boil the solution; carbonic acid gas escapes, and the carbonate is again precipitated.

This experiment will serve to show how chalk is kept in solution in ordinary well-waters, giving the property of "hardness," and how the *fur* or stone-like deposit in tea-kettles and boilers is formed. It should be here stated that sulphate of calcium produces similar hardness, and that these, with small quantities of the sulphate and carbonate of magnesium, constitute the hardening constituents of well-waters, a curd (oleate of calcium or magnesium) being formed whenever soap is used with such waters. An enormous amount of soap is wasted through the employment of hard water for washing-purposes. The hardness produced by the earthy carbonates is termed "temporary hardness," because removable by ebullition; that by the earthy sulphates "permanent hardness," because unaffected by ebullition. The addition of lime-water or a mixture of lime and water removes temporary hardness (react. 2, page 366), and carbonate of sodium, "washing-soda," both temporary and permanent hardness, in the latter case sulphate of sodium remaining in solution. Carbonate of barium (ground witherite) also decomposes sulphates of calcium and magnesium, sulphate of barium being precipitated and carbonates of calcium or magnesium formed; the latter and the carbonates originally in the water may then be precipitated by ebullition or by the action of lime-water. But the injurious effects of barium salts on man

and the lower animals prevents the carbonate being used for purifying water for drinking-purposes, as by accident or an unforeseen reaction a portion might become dissolved.

QUESTIONS AND EXERCISES.

523. Name the chief natural carbonates.
524. What are the formulæ of carbonic acid and carbonic acid gas?
525. Adduce evidence of the existence of true carbonic acid.
526. Trace the steps by which the carbonic constituent of chalk is transferred to sodium by the process usually adopted in alkali-works—the manufacture of “soda.”
527. Carbonic acid gas is constantly exhaled from the lungs of animals; why does it not accumulate in the atmosphere?
528. What is the effect of pressure on carbonic acid gas?
529. State the specific gravity of carbonic acid gas.
530. By what processes may carbonic acid gas be obtained for experimental and manufacturing purposes?
531. Describe the action of carbonic acid gas on the carbonates of potassium or sodium.
532. How may carbonic acid be detected in expired air?
533. To what extent is carbonic acid gas heavier than air?
534. Work sums showing what quantity of chalk (90 per cent. pure) will be required to furnish the carbonic acid necessary to convert one ton of carbonate of potassium (containing 83 per cent. of K_2CO_3) into acid carbonate, supposing no gas to be wasted.—*Ans.* 1500 lbs.
535. Define “hardness” in water.
536. How may the presence of carbonates be demonstrated?

OXALIC ACID AND OTHER OXALATES.

Formula of Oxalic Acid $H_2C_2O_4$, $2H_2O$.
Molecular weight 126.

Sources.—Oxalates occur in nature in the juices of some plants, as wood-sorrel, rhubarb, the common dock, and certain lichens; but the hydrogen oxalate (oxalic acid) and other oxalates are all made artificially. The carbon of many organic

substances yields oxalic acid when those substances are boiled with nitric acid, and an alkaline oxalate when they are roasted with a mixture of the hydrates of potassium and sodium.

Experimental process.—On the small scale, a mixture of nitric acid 10 parts, loaf-sugar 2 parts, and water 3 parts, quickly yields the acid. Abundance of red fumes are at first evolved. On cooling, crystals are deposited. A more dilute acid, kept warm, acts more slowly but yields a larger product.

Manufacturing process.—On the large scale, sawdust is roasted with alkalis, resulting oxalate of sodium decomposed by lime with formation of oxalate of calcium, the latter digested with sulphuric acid, and the liberated oxalic acid (Oxalic Acid of Commerce, B. P.) made commercially pure by recrystallization (Oxalic Acid, Purified, B. P.).

Chemically pure oxalic acid.—The acid made from sugar, recrystallized two or three times, is quite pure. Commercial acid should be mixed with insufficient water for complete solution, and the mixture occasionally shaken. Impurities remain undissolved, and the saturated aqueous solution evaporated yields crystals which seldom require to be recrystallized.

Quantivalence.—The elements represented by the formula C_2O_4 are those characteristic of oxalates. They form a bivalent grouping; hence normal oxalates ($R'_2C_2O_4$) and acid oxalates ($R'HC_2O_4$) exist.

Salt of sorrel is a crystalline compound of oxalic acid with acid potassium oxalate, the crystals containing two molecules of water of crystallization ($KHC_2O_4, H_2G_2O_4, 2H_2O$).

Analytical Reactions (Tests).

First Analytical Reaction.—To solution of an oxalate (oxalate of ammonium, *e.g.*) add solution of chloride of calcium; a white precipitate falls. Add to the precipitate excess of acetic acid; it is insoluble. Add hydrochloric acid; the precipitate is dissolved.

The formation of a white precipitate on adding a calcium or barium salt, insoluble in acetic but soluble in hydrochloric or nitric acid, is usually sufficient proof of the presence of an oxalate. The action of the liquid on litmus paper, effervescence with carbonate of sodium, and absence of metals, would indicate that the oxalate is that of hydrogen, oxalic acid.

Note.—The barium oxalate is slightly soluble in acetic acid (Souhay and Lenssen), and enough may be dissolved by this acid from a mixed barium precipitate (produced on adding chloride or nitrate of barium to a solution of mixed salts) to give the foregoing reaction on adding chloride of calcium to the filtered acetic liquid—an effect sometimes useful in the analysis of mixed substances (Davies).

Antidote.—In cases of poisoning by oxalic acid or salt of sorrel, chalk and water may be administered as a chemical antidote (with the view of producing the insoluble oxalate of calcium), emetics and the stomach-pump being used as soon as possible.

Second Analytical Reaction.—Heat a fragment of any dry common fixed metallic oxalate (an oxalate of potassium for example) in a test-tube; decomposition occurs, carbonic oxide (CO) (a gas that will be noticed subsequently) is liberated, and a carbonate of the metal remains. Add water and then an acid to the residue; effervescence occurs.

This is a ready test for ordinary insoluble oxalates, and is trustworthy if, on heating the substance, no charring occurs, or not more than gives a grey colour to the residue. Organic salts of metals decompose when heated, and leave a residue of carbonate, but, except in the case of oxalates, the residue is always accompanied by much charcoal. Insoluble oxalates and organic salts of such metals as lead and silver are, of course, liable to be reduced to oxide or even metal by heat. Such oxalates may be decomposed by boiling with solution of carbonate of sodium, filtering, and testing the filtrate for oxalates by the chloride of calcium test.

Other Analytical Reactions.—Nitrate of silver gives, with oxalates, white oxalate of silver ($\text{Ag}_2\text{C}_2\text{O}_4$).—Dry oxalates are decomposed when heated with strong sulphuric acid, carbonic oxide and carbonic acid gases escaping. If much of the substance be operated on, the gas may be washed with an alkali, the carbonic acid be thus removed, and the carbonic oxide be ignited; it will be found to burn with a characteristic bluish flame.—Oxalates, when mixed with water, black oxide of manganese (free from carbonates), and sulphuric acid,

yield carbonic acid gas, which may be tested by lime-water in the usual manner.—Not only such insoluble oxalates as those of lead and silver above referred to, but any common insoluble oxalate, such as that of calcium or magnesium, may be decomposed by ebullition with solution of carbonate of sodium; after filtration the oxalic radical will be found in the clear liquid as soluble oxalate of sodium.

Test of Purity.—"Purified Oxalic Acid . . . is entirely dissipated by a heat below 450° F." (B. P.).

QUESTIONS AND EXERCISES.

537. Explain the constitution of oxalates.
538. State how oxalates are obtained.
539. What is the quantivalence of the oxalic radical?
540. Give the formula of "salt of sorrel."
541. Mention the chief test for oxalic acid and other soluble oxalates.
542. Name the antidote for oxalic acid, and describe its action.
543. By what reactions are insoluble oxalates recognised?

TARTARIC ACID AND OTHER TARTRATES.

Formula of Tartaric Acid $\text{H}_2\text{C}_4\text{H}_4\text{O}_6$, or H_2T .
Molecular weight 150.

Sources.—Tartrates exist in the juice of many fruits; but it is from that of the grape that our supplies are usually obtained. Grape-juice contains much acid tartrate of potassium, which is gradually deposited when the juice is fermented, as in making wine; for acid tartrate of potassium, not very soluble in aqueous liquids, is still less so in spirituous, and hence crystallizes out as the sugar of the grape-juice is gradually converted into alcohol. It is found, with tartrate of calcium, lining the vessels in which wine is kept; and it is from this crude substance, termed *argal* or *argol*, also from

the albumenoid yeasty matter or "lees" deposited at the same time, as well as from what tartrate may be remaining in the marc left after the juice has been pressed from the grapes, that by rough recrystallization "tartar," containing 6 or 7 per cent. or more of tartrate of calcium, is obtained. From the tartar, tartaric acid and other tartrates are prepared. In old dried grapes (Raisins; *Uvæ*, B. P.) crystalline masses of tartar and of grape-sugar are constantly met with.

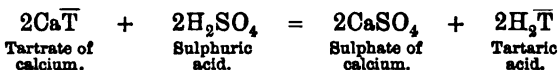
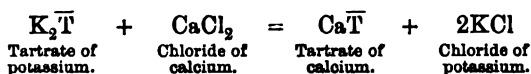
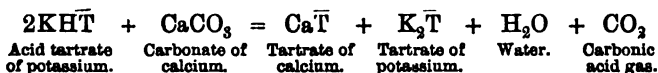
Cream of tartar purified by crystallization (*Potassæ Tartras Acida*, B. P.) occurs as "a gritty white powder, or fragments of cakes crystallized on one surface;" of a pleasant acid taste, soluble in 180 parts of cold and 6 of boiling water, insoluble in spirit.*

Quantivalence.—The elements represented by the formula $C_4H_4O_6$ are those characteristic of tartrates. They form a bivalent grouping; hence normal tartrates ($R'_2\bar{T}$) and acid tartrates ($R'H\bar{T}$) exist. Tartrate of potassium, the *Potassæ Tartras* of the British Pharmacopœia ($K_2C_4H_4O_6$), and Rochelle Salt, or tartrate of potassium and sodium, the official *Soda Tartarata*, are illustrations of normal tartrates, while Cream of Tartar, $KHC_4H_4O_6$, is an example of acid tartrates. The only official tartrate not apparently included in these general formulæ is tartar-emetic (*Antimonium Tartaratum*, B. P.), which is sometimes regarded as the double tartrate of potassium and a hypothetical radical, antimonyl (SbO), thus, $KSbOC_4H_4O_6$. Probably, however, it is but an oxytartrate of antimony ($Sb_2O_2\bar{T}$) with normal tartrate of potassium ($K_2\bar{T}$); for there are several oxycompounds of antimony analogous to the oxycompounds of bismuth that have been described (p. 290), normal salts partially decomposed by water into oxides, and many of these oxycompounds readily unite with normal salts of other basylous radicals. Tartar-emetic would thus be oxytartrate of antimony with tartrate of potassium [$Sb_2O_2\bar{T}$, $K_2\bar{T}$, or $Sb_2O_2C_4H_4O_6$, $K_2C_4H_4O_6$, or $K'Sb'''(C_4H_4O_6)''O''$].

* A boiling solution of tartar yields a floating crust of minute crystals on cooling—just as milk yields a floating layer of cream; hence the term *cream of tartar*. "It is called *tartar*," says Paracelsus, "because it produces oil, water, tincture and salt, which burn the patient as tartarus does." *Tartarus* is Latin (*Tátrapos*, Tartaros, Greek) for *hell*. The products of its destructive distillation are certainly somewhat irritating in taste and smell; and the "salt" (carbonate of potassium) that is left is diuretic and in larger quantities powerfully corrosive.

Tartaric Acid.

Tartaric Acid (*Acidum Tartaricum*, B. P.) is obtained, according to the British Pharmacopœia, by boiling cream of tartar (*Potassæ Tartras Acida*, B. P.) with water, adding chalk till effervescence ceases, and then chloride of calcium so long as a precipitate falls; the two portions of tartrate of calcium thus consecutively formed are thoroughly washed, treated with sulphuric acid, the mixture boiled for a short time, resulting sulphate of calcium mostly separated by filtration, the filtrate concentrated by evaporation, any sulphate of calcium that may have deposited removed as before, and concentration continued until the solution is strong enough to crystallize. Tartrate of calcium from nine ounces of cream of tartar requires five ounces by weight of sulphuric acid for complete decomposition.



Tartaric acid occurs in trade in colourless crystals, or the same powdered. It is strongly acid and readily soluble in water or spirit. One part in eight of water and two of spirit of wine forms "Solution of Tartaric Acid," B. P. Its aqueous solution is not stable.

Parcels of tartaric acid often contain crystals of an allotropic or physically isomeric modification (*vide* "Allotropy" and "Isomerism" in Index). It is termed *Paratartaric acid* (*παρά, para*, beside) or *Racemic acid* (*racemus*, a bunch of grapes), and is a combination of ordinary tartaric acid, whose solution twists a ray of polarized light to the right hand (dextrotartaric or dextroracemic acid), and of lævotartaric or lævoracemic acid, whose solution twists a polarized ray to the left. Racemic acid is inactive in this respect, the opposite properties of its constituents neutralizing each other. Racemic acid is less soluble in alcohol than tartaric acid.

REACTIONS.

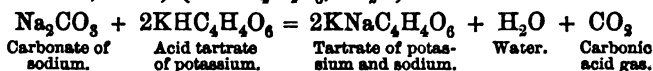
Tartrate of Potassium.

Synthetical Reactions.—To a small quantity of a strong solution of carbonate of potassium add acid tartrate of potassium, so long as effervescence occurs; the resulting liquid is solution of normal tartrate of potassium (*Potassæ Tartras*, B. P.) ($K_2C_4H_4O_6$), crystals of which may be obtained on evaporation.

Note.—This is a common method of converting an acid salt of a bivalent acidulous radical into a normal salt. The carbonate added need not be a carbonate of the same, but may be of a different metal; compounds like Rochelle salt ($KNaC_4H_4O_6$) are then obtained. Thus:—

Tartrate of Potassium and Sodium.

To a strong hot solution of carbonate of sodium add acid tartrate of potassium until effervescence ceases; the resulting liquid is solution of tartrate of potassium and sodium: on cooling, it yields crystals. This is the official process (*Soda Tartarata*, B. P.) ($KNaC_4H_4O_6, 4H_2O$).



Crystals of Rochelle salt are usually halves of colourless transparent right rhombic prisms, slightly efflorescent in dry air, soluble in five parts of boiling water. Tartrate of potassium is slightly deliquescent, soluble in about four parts of boiling water.

Equivalent Weights of Tartaric Acid, Carbonate of Potassium, Bicarbonate of Potassium, Carbonate of Sodium (crystallized), Bicarbonate of Sodium, and Carbonates of Ammonium and Magnesium: repeated for 20 parts of each (and, incidentally, for other proportions).

Tart. Acid . .	$H_2C_4H_4O_6$	=150	20	18½	15	10½	17½	25½	31½
Carb. Potas. .	K_2CO_3 (of 84 per cent.) . .	=164	22	20	16½	11½	19½	28	34½
Bicarb. Pot. .	$2(KHCO_3)$	=200	26½	24½	20	14	23½	34	42
Carb. Sod. .	$Na_2CO_3, 10H_2O$	=286	38	34½	28½	20	34	49½	60
Bicarb. Sod. .	$2(NaHCO_3)$	=168	22½	20½	16½	11½	20	28½	35½
Carb. Ammon.	$(N_2H_4C_2O_4) + 2$	=118	15½	14½	11½	8½	14	20	24½
Carb. Magnes.	$(MgCO_3), Mg_2HO, 4H_2O + 4$ =95.5		12½	11½	9½	6½	11½	16	20

Thus 20 parts (grains or other weights) of tartaric acid neutralize 22 of carbonate of potassium, $26\frac{1}{2}$ of bicarbonate of potassium, 38 of carbonate of sodium, $22\frac{1}{2}$ of bicarbonate of sodium, $15\frac{1}{2}$ of carbonate of ammonium, or $12\frac{1}{2}$ of carbonate of magnesium. Other quantities of tartaric acid ($18\frac{1}{2}$, 15, $10\frac{1}{2}$, $17\frac{1}{2}$, $25\frac{1}{2}$, $31\frac{1}{2}$) saturate the amounts of salts mentioned in the other columns and *vice versâ*. A similar Table for Citric Acid will be found at p. 380, and for both acids in the Appendix. These Tables afford good illustrations of some of the laws of chemical combination (pp. 43, 222). The reader should verify a few of the numbers by calculation from the atomic weights of the elements concerned in the reactions, remembering that the salts formed are considered to be neutral in constitution. In medical practice effervescing saline draughts are often designedly prescribed to contain an amount of acid or alkali considerably in excess of the proportions required for perfect neutrality.

A common form of *Seidlitz Powder* consists of 3 parts of Rochelle salt (120 grains) with 1 (40 grains) of acid carbonate of sodium (the mixture usually wrapped in blue paper) and 1 (40 grains) of tartaric acid (wrapped in white paper). When administered, the latter is dissolved in a tumbler rather more than half full of water, the former added, and the mixture drank during effervescence. It will be seen that the salts swallowed are tartrate of potassium and sodium (KNaT , $4\text{H}_2\text{O}$), tartrate of sodium (Na_2T , $2\text{H}_2\text{O}$), and acid tartrate of sodium or of potassium. The last-mentioned salt results because $11\frac{1}{2}$ per cent. ($4\frac{1}{2}$ grains) of the tartaric acid is in excess of the quantity necessary for the formation of neutral tartrate of sodium. This amount of acid salt gives agreeable acidity to the draught. The American formula (*Pulveres Effervescentes Aperientes*, U. S. P.) includes rather less tartaric acid, so that only neutral salts are formed.

Analytical Reactions (Tests).

First Analytical Reaction.—To solution of any normal tartrate, or tartaric acid made neutral by solution of soda, add solution of chloride of calcium; a white precipitate, tartrate of calcium, falls. Collect the precipitate on a filter, wash, place a small quantity in a test-tube, and add solution of potash; on stirring the mixture the precipitate dissolves.

Heat the solution; the tartrate of calcium is again precipitated.

In the above reaction a fair amount of the chloride of calcium solution should be added at once, and the whole test performed without delay, or the calcium tartrate will assume a crystalline character, and be with difficulty dissolved by the potash.

The solubility of tartrate of calcium in cold potash solution enables the analyst to distinguish between tartrates and citrates, otherwise a difficult matter. Citrate of calcium is not soluble, or only to a very slight extent, in the alkali. The absence of much ammoniacal salt must be insured, citrate, as well as tartrate, of calcium, being soluble in solutions of salts of ammonium.

Second Analytical Reaction.—Acidulate a solution of a tartrate with acetic acid, add acetate of potassium, and well stir the mixture; a crystalline precipitate of acid tartrate of potassium slowly separates.

This reaction is not applicable in testing for very small quantities of tartrates, the acid tartrate of potassium being not altogether insoluble. The precipitate being insoluble in alcohol, however, the addition of spirit of wine renders the test far more delicate.

Third Analytical Reaction.—To a neutral solution of a tartrate add solution of nitrate of silver; a white precipitate of tartrate of silver, $\text{Ag}_2\text{C}_4\text{H}_4\text{O}_6$, falls. Boil the mixture; it blackens, owing to the reduction of the salt to metallic silver. Or, before boiling, add a drop, or less, of ammonia; a mirror will form on the tube—adhering well to the glass if the tube was thoroughly cleansed. Even an insoluble tartrate, placed in a dry tube with a few fragments of nitrate of silver and a drop, or less, of ammonia added, gives a mirror-like character to each fragment of the silver salt when the tube is gently rotated some inches above a flame.

Fourth Analytical Reaction.—To a neutral or alkaline solution of a tartrate add a few drops of solution of permanganate of potassium, and slowly heat the test-tube; the colour is

discharged, an oxide of manganese being precipitated. Citrates only reduce the permanganate to green manganate.

Other Reactions.—Tartrates heated with strong sulphuric acid char immediately.—Tartaric acid and the soluble tartrates prevent the precipitation of ferric and other hydrates by alkalies, soluble double tartrates being formed (which on evaporation yield liquids that do not crystallize, but, spread on sheets of glass, dry up to thin transparent plates or scales). The potassio-tartrate of iron (*Ferrum Tartaratum*, B. P.) is a preparation of this kind.—Tartrates decompose when heated, carbonates being formed and carbon set free, the gaseous products having a peculiar, more or less characteristic smell, resembling that of burnt sugar.

QUESTIONS AND EXERCISES.

544. State the origin of tartaric acid and other tartrates, and explain the deposition of argol, crude acid tartrate of potassium, during the manufacture of wine.

545. What is the chemical formula and what are the characters of "cream of tartar"?

546. Mention the formula and quantivalence of the tartaric radical.

547. Write formulæ of normal, acid, and double tartrates, tartar-emetic being treated as an oxytartrate of antimony with tartrate of potassium.

548. Give equations or diagrams illustrative of the production of tartaric acid from cream of tartar.

549. By what general process may normal or double tartrates be obtained from acid tartrate of potassium?

550. Work out sums proving the correctness of some of the figures given on p. 374 as showing the saturating-power of tartaric acid for various quantities of different carbonates, and give diagrams or equations of the reactions.

551. State the names and work sums showing quantities of the salts resulting from the admixture of 120 grains of tartrate of potassium and sodium, 40 grains of acid carbonate of sodium, and 40 grains of tartaric acid (seidlitz powder).

552. Enumerate the tests for tartrates, and explain the effects of heat on tartrates of the metals.

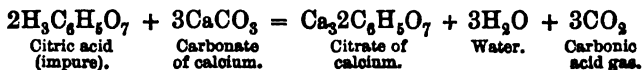
CITRIC ACID AND OTHER CITRATES.

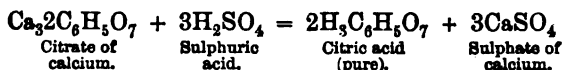
Formula of Citric Acid $\text{H}_3\text{C}_6\text{H}_5\text{O}_7$, H_2O or $\text{H}_3\bar{\text{C}}\text{i}$, Aq.

Molecular weight 210.

Source.—Citric acid (*Acidum Citricum*, B. P.) exists in the juice of many of our common garden fruits. The pulp of the fruit of *Tamarindus indica* (*Tamarindus*, B. P.) contains nearly 10 per cent. (in addition to 1.5 of tartaric acid, .5 of malic acid, and 3 per cent. of acid tartrate of potassium). But it is from the lemon or lime that the acid of commerce is usually obtained. For this purpose concentrated lemon-juice is exported from Sicily, concentrated bergamot juice from the Calabrian coast of South Italy, and concentrated lime-juice from the West Indies. The lime-fruit from *Citrus bergamia* is official in the Pharmacopœia of India.

Process.—The British Pharmacopœia directs that the hot lemon-juice (4 pints) be saturated by powdered chalk, that is, whiting ($4\frac{1}{2}$ ounces), the resulting citrate of calcium collected on a filter, washed with hot water till the liquor passes from it colourless (by which not only the colouring-matter but the mucilage, sugar, and other constituents of the juice are got rid of), then mixed with cold water (1 pint), decomposed by sulphuric acid ($2\frac{1}{2}$ fluid ounces in $1\frac{1}{2}$ pint of water), the mixture boiled for half an hour, filtered, the solution evaporated to a density of 1.21, set aside for 24 hours, then poured off from any deposit of crystalline sulphate of calcium, further concentrated and set aside to crystallize. If the quantity of citrate of calcium to be decomposed is indefinite, the sulphuric acid may be added until a little of the supernatant fluid gives, after a minute or two, a precipitate with solution of chloride of calcium. The concentrated citric solution generally crystallizes very slowly. Shaken violently, however, in a bottle with a granule or two of solid acid, it quickly yields its citric acid in a pulverulent form, and this drained and redissolved in a very small quantity of hot water yields crystals fairly quickly (Warington).





Quantivalence.—The elements represented by the formula $\text{C}_6\text{H}_5\text{O}_7$ are those characteristic of citrates. They form a trivalent grouping; hence three classes of salts may exist—one, two or three atoms of the basylous hydrogen in the acid, $\text{H}_3\text{C}_6\text{H}_5\text{O}_7$, being displaced by equivalent proportions of other basylous radicals.

Citric acid itself is the only citric compound of much direct importance to the pharmacist. It usually occurs in colourless crystals soluble in half their weight of boiling and three-fourths of cold water, less soluble in spirit, and insoluble in ether. A solution of about 34 grains in 1 ounce of water forms a sort of artificial lemon-juice. Citrates heated with strong sulphuric acid to about 212°F . evolve carbonic oxide gas, and at higher temperatures acetone and carbonic acid gas.

The artificial production of citric acid has been accomplished by Grimaux and Adam, who starting with glycerine, produce certain chloro- and cyano-derivatives and ultimately citric acid itself.

Action of heat on citric acid.—Citric acid slowly heated first loses its water of crystallization; afterwards (347°F .) the elements of another molecule of water are evolved and a residue obtained from which ether extracts *aconitic acid* ($\text{H}_3\text{C}_6\text{H}_5\text{O}_6$), identical with the aconitic acid (and the acid first termed *equisetie*) in various species of *Aconitum* and *Equisetum*.

The official Lemon-juice (*Succus Limonum*, B. P.) is to be freshly expressed from the ripe fruit, and contain an average of 32.5 grains of citric acid in 1 fluid ounce. The acidity may be ascertained by adding solution of potash or soda (the strength of which has been previously determined with pure crystals of citric acid) till red litmus-paper is fairly turned blue. Before applying this test to commercial specimens, the absence of notable quantities of sulphuric, hydrochloric, acetic, tartaric, or other acid must be insured by application of appropriate re-agents. (See also "Lemon-juice" in Index.)

Equivalent Weights of Citric Acid, Carbonate of Potassium, Bicarbonate of Potassium, Carbonate of Sodium (crystal-

lized), Bicarbonate of Sodium, and Carbonates of Ammonium and Magnesium: repeated for 20 parts of each (and, incidentally, for other proportions).

Citric Acid	$\text{H}_3\text{C}_6\text{H}_5\text{O}_7, \text{H}_2\text{O}$	$\dots = 210$	20	17	14	9	16	23	29
Carb. Pot.	$(\text{K}_2\text{CO}_3; \text{ of } 8\frac{1}{2} \text{ per cent.}) + 2 \times 3$	$= 240$	23	20	16	11	19	28	34
Bicarb. Pot.	$3(\text{KHCO}_3)$	$\dots = 300$	28	24	20	14	23	34	41
Carb. Sod.	$(\text{Na}_2\text{CO}_3, 10\text{H}_2\text{O}) + 2 \times 3$	$= 420$	40	34	28	20	34	49	60
Bicarb. Sod.	$3(\text{NaHCO}_3)$	$\dots = 252$	24	20	16	11	20	28	35
Carb. Amm.	$(\text{N}_2\text{H}_4\text{C}_2\text{O}_6) + 4 \times 3$	$= 177$	16	14	11	8	14	20	24
Carb. Mag.	$(\text{MgCO}_3, \text{Mg}_2\text{HO}, 4\text{H}_2\text{O} + 8 \times 3 = 144)$	$= 144$	13	11	9	6	11	16	20

Thus 20 parts (grains, or other weights) of citric acid neutralize $23\frac{1}{2}$ of carbonate of potassium, $28\frac{1}{2}$ of bicarbonate of potassium, 40 of carbonate of sodium, $24\frac{1}{2}$ of bicarbonate of sodium, $16\frac{1}{2}$ of carbonate of ammonium, or $13\frac{1}{2}$ of carbonate of magnesium. Other quantities of acid (17, 14, $9\frac{1}{2}$, $16\frac{1}{2}$, $23\frac{1}{2}$, $29\frac{1}{2}$) saturate the amounts of salts mentioned in the other columns and *vice versa*.

This Table, the similar one for tartaric acid (p. 374), and that for both acids (*vide* Appendix) afford good illustrations of some of the laws of chemical combination (pp. 43, 222). The reader should verify a few of the numbers by calculation from the atomic weights of the elements concerned in the reactions, remembering that the salts formed are considered to be neutral in constitution. In medical practice, effervescing saline draughts are often designedly prescribed to contain an amount of acid or alkali considerably in excess of the proportions required for perfect neutrality.

Analytical Reactions (Tests).

First Analytical Reaction.—To a dilute solution of any neutral citrate, or citric acid carefully neutralized by alkali, add solution of chloride of calcium and boil; a white precipitate, citrate of calcium ($\text{Ca}_3\text{C}_6\text{H}_5\text{O}_7$), falls. Treat the precipitate as for tartrate of calcium (p. 375); it is not perceptibly dissolved by the potash.

A mixture of citrates and tartrates can be separated by this reaction. They are precipitated as calcium salts, and the rapidly washed precipitate mixed with solution of potash, diluted and filtered; the filtrate contains the tartrate, which is shown to be present by reprecipitation on boiling. The pre-

precipitate still on the filter is washed, dissolved in solution of chloride of ammonium, and the solution boiled; the citrate of calcium is reprecipitated. The presence of much sugar interferes with this reaction. A dilute solution of a citrate is not precipitated by chloride of calcium until the liquid is heated: precipitation from a strong solution, also, is not thoroughly complete without ebullition of the mixture. This reaction is not thoroughly satisfactory, citrate of calcium being *slightly* soluble in alkalies, in the solutions of salts produced in the reaction, and, to a very slight extent, even in cold water.

Second Analytical Reaction.—To a neutral solution of a citrate add solution of nitrate of silver; a white precipitate of citrate of silver (Ag_3Ci) falls. Boil the mixture; the precipitate does not blacken as tartrate of silver does, or only after long boiling.

Third Analytical Reaction.—To a neutral or alkaline solution of a citrate add a few drops of solution of permanganate of potassium and slowly heat the test-tube; reduction to manganate only occurs, a green or reddish-green solution resulting. Tartrates reduce the permanganate entirely.

Other Analytical Reactions.—Citric acid forms no precipitate corresponding with the acid tartrate of potassium.—Lime-water, in excess, gives no precipitate with citric acid or citrates, unless the solution is boiled, citrate of calcium being slightly soluble in cold but not in hot water; it usually precipitates tartrates in the cold.—Citrates, when heated with strong sulphuric acid, do not char immediately.—Citric acid and citrates prevent the precipitation of oxide of iron by alkalies, soluble double compounds being formed. The *Ferri et Ammoniac Citras*, B. P., is a preparation of this kind.—Metallic citrates decompose when heated, carbonates being formed and carbon set free; the odour of the gaseous products is not so characteristic as that of tartrates.—According to Cailliet a cold saturated solution of red chromate of potassium turns a solution of tartaric acid dark brown, carbonic acid gas being evolved, while a solution of citric acid only slowly becomes of a light brown.

QUESTIONS AND EXERCISES.

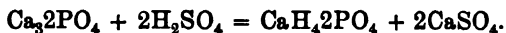
553. What is the source of citric acid ?
 554. Describe the method by which citric acid is prepared, giving diagrams.
 555. Illustrate by formulæ the various classes of tartrates and citrates.
 556. State the average proportion of citric acid in lemon-juice.
 557. Work out sums proving the correctness of some of the figures given on page 380 as showing the saturating power of citric acid for various carbonates.
 558. What are the tests for citrates ?
 559. How are tartrates separated from citrates ?
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PHOSPHORIC ACID AND OTHER PHOSPHATES.

Formula of Phosphoric Acid H_3PO_4 . Molecular weight 98.

Source.—The source of the ordinary normal phosphates and of phosphorus itself (*Phosphorus*, B. P.) is the normal phosphate of calcium (Ca_3PO_4). It is the chief constituent of the bones of animals, being derived from the plants on which they feed, plants again obtaining it from the soil. Compounds of phosphorus are also met with in the brain, nerves, muscles, blood, saliva, and, according to Kirkes, even in tissues so simple that one must assume that the compounds are necessary constituents of the substance of the primary cell. They escape from the system both in the urine and in the fæces.

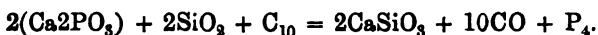
Process.—Phosphorus ($\text{P}=31$) is obtained from bones by the following processes:—The bones are burnt to remove all traces of animal matter. The resulting *bone-earth* is treated with sulphuric acid and water, by which an acid phosphate of calcium (CaH_2PO_4), often called *superphosphate of lime*, is produced:—



The acid phosphate (strained from the sulphate and evaporated to dryness) is mixed with charcoal and sand and heated to dull redness in an iron pot. At this stage water escapes and metaphosphate of calcium (Ca_2PO_3 , *see* Index) remains:—



The mixture is then transferred to a retort and distilled at a strong red heat; a silicate of calcium (CaSiO_3) is formed and remains in the retort, phosphorus vapour is evolved and condensed under water, and carbonic oxide gas escapes:—



It is purified by melting under water containing sulphuric acid and red chromate of potassium.

Properties.—Phosphorus is “a semitransparent, colourless, wax-like solid (in sticks or cakes), which emits white vapours when exposed to the air. Specific gravity 1.77. It is soft and flexible at common temperatures, melts at 110°F ., ignites in the air at a temperature a little above its melting-point, burning with a luminous flame and producing dense white fumes. It is very poisonous. Insoluble in water, but soluble in ether and in boiling oil of turpentine,” also in bisulphide of carbon. It is soluble in oil which has been previously heated for a short time to about 300°F . to expel moisture—12 grains in 4 fluid ounces of dried almond-oil constituting Phosphorated Oil, *Oleum Phosphoratum*, B. P. A mixture, or rather a solution, of phosphorus in balsam of tolu and yellow wax, forms the official Phosphorus Pill (*Pilula Phosphori*, B. P.).

Granulated or pulverulent phosphorus is obtained by placing a portion under equal parts of spirit and water in a bottle, standing the bottle in warm water till the phosphorus melts, then inserting the stopper (glass, not cork) and shaking the whole till cold.

Red or Amorphous Phosphorus.—Ordinary phosphorus kept at a temperature of about 450°Fahr ., in an atmosphere from which air is excluded, becomes red, opaque, insoluble in liquids in which ordinary phosphorus is soluble, oxidizes extremely slowly, and only ignites when heated to near 500°Fahr . It is used in the manufacture of several varieties of lucifer-matches, not emitting the poisonous jaw-destroying fumes given by ordinary phosphorus.

Quantivalence.—The atom of phosphorus is quinquivalent, as seen in the pentachloride (PCl_5) and oxychloride (PCl_3O); but it often exhibits trivalent activity, as seen in the trichloride (PCl_3) and trihydride (PH_3).

Molecular weight.—Phosphorus is an exception to the rule that the atomic weights (in grains, grammes, etc.) of elements occupy similar volumes of vapour at similar tempera-

tures, the equivalent weight of phosphorus (31) only giving half such a volume. Hence while the molecular weights, that is, double the atomic weights, of oxygen ($O_2=32$), hydrogen ($H_2=2$), nitrogen ($N_2=28$), etc. give a similar bulk of vapour at any given temperature, the double atomic weight of phosphorus ($P_2=62$) only gives half this bulk; that is, four times the atomic weight of phosphorus must be taken to obtain the whole bulk. It would appear therefore that the molecule of phosphorus contains four atoms ($P_4=124$). As with sulphur, however, phosphorus in the state ordinarily known to us may be abnormal, and a variety yet be found in which the molecular weight is double the atomic weight.

Phosphoric Acid.

The chief use of phosphorus in pharmacy is in the formation of Diluted Phosphoric Acid. Phosphorus is boiled with nitric acid and water until dissolved. The solution, evaporated to a low bulk to remove nitrous compounds, and rediluted so as to contain nearly 14 (13·8) per cent. of acid (H_3PO_4), equivalent to 10 per cent. of phosphoric anhydride (P_2O_5), constitutes the *Acidum Phosphoricum Dilutum*, B. P., a colourless sour liquid of specific gravity 1·08. If the necessary appliances are at hand, four or five ounces of this acid may be prepared by the official process as follows:—A quarter of a pint is made by boiling together, in a retort attached to a Liebig's condenser, 103 grains of phosphorus, $1\frac{1}{2}$ fluid ounce of the official nitric acid, and 2 ounces of water. When about 1 oz. of water has distilled over it should be returned to the retort, and the operation repeated until the phosphorus has disappeared.



The liquid remaining in the retort is then transferred to a dish (preferably of platinum), evaporated down to about half an ounce, and, lastly, diluted with distilled water to 5 fluid ounces.

The use of the water in this process is to moderate the

reaction. Strong hot nitric acid oxidizes phosphorus with almost explosive rapidity, hence the acid must be diluted in the first instance, and be rediluted from time to time to prevent its becoming too strong by loss of water. Time is saved by using a strong acid, but in that case constant supervision is necessary in order that water may be added, or the temperature otherwise reduced, when the action becomes too violent. Deficiency of nitric acid must also be avoided, or some phosphorus acid (H_2PHO_3) will be formed.

Markoe, also to economise time, modifies the process by adding for every ounce of phosphorus 4 or 5 grains of iodine, and, drop by drop, 25 or 30 drops of bromine. The iodine and bromine unite with the phosphorus with a readiness or even violence that would be explosive if not controlled by the presence of the cold fluids,—further cooled, if necessary, by immersing the vessel in cold water. Iodide of phosphorus (PI_3) and bromide of phosphorus (PBr_3) are at once formed. These in the presence of water immediately yield hydriodic and hydrobromic acids (HI , HBr) and phosphoric acid. The nitric acid attacks the hydriodic and hydrobromic acids, forming the lower oxides of nitrogen, which escape as gas, water, and free iodine and bromine. The latter unite with more phosphorus, and the reactions are repeated. This carrying power of a little iodine or bromine or both would perhaps be indefinitely prolonged if no vapour of these elements or their acids escaped with the gases. The phosphorus having disappeared, excess of nitric acid is got rid of roughly by dropping in clean rags or paper (nitric oxide, carbonic acid gas, and water being formed), and, the last portions, by adding oxalic acid (which even still more readily yields similar products). Evaporation to a syrupy consistence finally removes all traces of iodine, bromine, oxalic acid, and moisture. The product is then diluted to any required extent.

Experimental process.—A flask, in the neck of which a funnel is inserted and a second funnel inverted, so that its mouth rests within the mouth of the first, is an efficient and convenient arrangement of apparatus for this process, especially if the operation be conducted slowly. (See fig. 40.)

Solution of phosphoric acid evaporated leaves a residue which melts at a low red heat, yielding *pyrophosphoric acid*, and, finally, *metaphosphoric acid* (*Glacial Phosphoric Acid*).

Phosphoric acid is also easily made from amorphous phosphorus (Mattison).

Fig. 40.



Prepared from bones phosphoric acid is apt to develop fungoid deposits (Jensen).

Quantivalence.—The elements represented by the formula PO_4 are those characteristic of phosphates. The grouping is trivalent; hence there may exist trimetallic or normal phosphates ($\text{M}'_3\text{PO}_4$), dimetallic acid phosphates ($\text{M}'_2\text{HPO}_4$), monometallic acid phosphates ($\text{M}'\text{H}_2\text{PO}_4$), and, lastly, trihydric phosphate (H_3PO_4), or common phosphoric acid. These are the ordinary phosphates, or *orthophosphates*, met with in nature or used in pharmacy; the rarer pyrophosphates and metaphosphates, as well as the phosphites and hypophosphites, will be mentioned subsequently.

Analytical Reactions (Tests).

First Analytical Reaction.—To an aqueous solution of a phosphate (e.g. Na_2HPO_4) add solution of sulphate of magnesium with which chloride of ammonium and ammonia have been mixed; a white crystalline precipitate of ammonio-magnesian phosphate falls (MgAmPO_4).

Chloride of ammonium is added to prevent the precipitation of hydrate of magnesium. Arseniates, which have close analogy to phosphates, give a precipitate of similar character with the magnesian reagent.

Second Analytical Reaction.—To a neutral aqueous solution of a phosphate add solution of nitrate of silver; light yellow

phosphate of silver (Ag_3PO_4) is precipitated. To a portion of the precipitate add ammonia; it dissolves. To another portion add nitric acid; it dissolves. By this reaction phosphates may be distinguished from their close allies the arseniates, arseniate of silver being of a chocolate colour.

Third Analytical Reaction.—To a solution (in a few drops of acid) of a phosphate insoluble in water (e.g. Ca_3PO_4) add the acetate of an alkali metal (easily made by adding to soda or ammonia in a test-tube excess of acetic acid), and then a drop or two of solution of perchloride of iron; yellowish white ferric phosphate (Fe_3PO_4) is precipitated. Too much of the ferric chloride must not be added, or ferric acetate will be produced, in which ferric phosphate is to some extent soluble.

To remove the whole of the phosphoric radical from the solution add ferric chloride so long as a precipitate is produced, and boil; ferric phosphate and oxyacetate are precipitated.

To obtain confirmatory evidence of the presence of phosphate in this precipitate and to separate the phosphoric radical as a pure unmixed phosphate, collect the precipitate on a filter, wash, drop some solution of ammonia on it, then sulphhydrate of ammonium, and finally wash with water; black ferrous sulphide remains on the filter, while phosphate of ammonium occurs in the filtrate. To the filtrate add a mixture of solutions of sulphate of magnesium and chloride of ammonium, and well stir; ammonio-magnesian phosphate is precipitated.

Fourth Analytical Reaction.—To diluted nitric acid add a little phosphate of calcium (or any other phosphate) and then solution of molybdate of ammonium, and gently heat; a yellow precipitate falls.

This precipitate contains what is termed phospho-molybdic acid, but is a compound of molybdic acid with phosphoric acid (about 4 per cent. of H_3PO_4) and ammonia (nearly 7 per cent.).

Molybdate of ammonium is obtained by roasting the native

sulphide of molybdenum (MoS_2) to molybdic oxide or anhydride (MoO_3), dissolving the latter in water, adding ammonia, evaporating, and crystallizing.

Molybdates having the following formulæ ($M = 1$ univalent atom of any metal) have been obtained:— M_2MoO_4 ; MHMoO_4 ; MHMoO_4 , H_2MoO_4 . According to Carrington, commercial molybdate of ammonium is commonly the intermediate of the three salts.

Note.—The foregoing two reactions are useful in the analysis of bone-earth, of other earthy phosphates, phosphate of iron, and all phosphates insoluble in water. Only arseniates give similar appearances; but the acid solution of these may be decomposed by agitation with sulphurous acid, ebullition, and subsequent treatment with sulphuretted hydrogen—arsenious sulphide, As_2S_3 , being then precipitated.

Other Analytical Reactions.—Solutions of barium and calcium salts give, with aqueous solutions of phosphates, white precipitates of the respective phosphates BaHPO_4 , or $\text{Ba}_2\text{P}_2\text{O}_7$, and CaHPO_4 , or $\text{Ca}_3\text{P}_2\text{O}_7$, all of which are soluble in acetic and the stronger acids.

QUESTIONS AND EXERCISES.

560. State the source of phosphorus.
561. Give equations or diagrams explanatory of the isolation of phosphorus from its natural compounds.
562. What is the composition of farmers' "superphosphate," and how is it prepared?
563. Enumerate the properties of phosphorus.
564. Mention some solvents of phosphorus.
565. How is the official Diluted Phosphoric Acid made?
566. Describe the precautions necessary to be observed in making this acid.
567. What is the strength of the official acid?
568. Write formulæ illustrative of all classes of orthophosphates.
569. Mention the chief tests for soluble and insoluble phosphates.
570. By what reactions may phosphates be distinguished from arseniates?

VANADIUM, V, 51.3, is a very rare element, and is here mentioned only because of its exceedingly interesting relationship to nitrogen, phosphorus, and arsenium. Discovered, but not isolated, by Sefström, and its compounds investigated by Berzelius, it has only of late years been obtained in the free state and fully studied by Roscoe.

$N_2O_5, N_2O_4, N_2O_3, N_2O_2, N_2O.$	$V_2O_5, V_2O_4, V_2O_3, V_2O_2, V_2O.$
Orthophosphates . R'_3PO_4	Orthovanadates . R'_3VO_4
Pyrophosphates . $R'_4P_2O_7$	Pyrovanadates . $R'_4V_2O_7$
Metaphosphates . $R'PO_3$	Metavanadates . $R'VO_3$

Isomorphous Minerals.

Apatite	$3(Ca_32PO_4), CaFl_2$
Pyromorphite	$3(Pb_32PO_4), PbCl_2$
Mimetesite	$3(Pb_32AsO_4), PbCl_2$
Vanadinite	$3(Pb_32VO_4), PbCl_2$

BORACIC ACID AND OTHER BORATES.

Formula of Boracic Acid H_3BO_3 . Molecular weight 62.

The composition of artificial boracic acid is expressed by the formula H_3BO_3 ; but at a temperature of 212° F. this body loses the elements of water and yields metaboracic acid, HBO_2 . The latter acid exists in the jets of steam (*fumerolles* or *suffioni*) that issue from the earth in some districts of Tuscany, and collects in the water of the *lagoni* (lagoons or little lakes) formed at the orifices of the steam-channels. This acid liquid, evaporated by aid of the waste natural steam and neutralized by carbonate of sodium, gives common borax ($2NaBO_3, 2HBO_3, 9H_2O$), possibly an acid metaborate of sodium with water of crystallization. It occurs "in transparent colourless crystals, sometimes slightly effloresced, with a weak alkaline reaction; insoluble in rectified spirit, soluble in water." Native borax, or *tincal*, and other borates, are also found in Thibet, Nevada, Peru, Chili, and, recently, in California in the Colorado district. The introduction of the natural borax from California has reduced the price to about one-half its former amount. The borax is represented as forming large portions of the crystalline bed of a dried-up lake. Fused borax readily dissolves metallic oxides, as will

have been already noticed in testing for cobalt and manganese. Hence, besides its use in medicine (*Borax*; *Mel Boracis*, B. P., and *Glycerinum Boracis*, B. P.), it is employed as a flux in refining and other metallurgic and ceramic operations.

Quantivalence.—The boracic radical is trivalent (BO_3'''); the metaboracic univalent (BO_2'); they have not been isolated.

The element boron, like carbon, occurs in the amorphous, graphitoidal, and crystalline conditions. It is a trivalent element (B'''), yielding definite salts, such as the chloride (BCl_3) and fluoride (BF_3). Its atomic weight is 11.

REACTIONS.

First Synthetical Reaction.—To a hot solution of a crystal of borax add a few drops of sulphuric acid and set aside; on cooling, crystalline scales of boracic acid (H_3BO_3) are deposited. They may be purified by collecting on a filter, slightly washing, drying, digesting in hot alcohol, filtering, and setting aside; pure boracic acid (B. P.) is deposited. The acid may also be recrystallized from water. Fifty grains dissolved in one ounce of rectified spirit constitutes "Solution of Boracic Acid," B. P.

When heated, these crystals lose water and yield, first, metaboracic acid (HBO_2), and subsequently, when fused, boracic anhydride (B_2O_3).

Boracic acid is a very weak compound. It only slowly decomposes carbonates, and resembles alkaline substances in colouring turmeric brown. Indeed the alkalinity of borax is as great as if it contained no acidulous radical. Boracic acid is said to be itself an antiseptic, but Endemann states that in preserving foods it acts by converting phosphates into acid phosphates and that the latter are the antiseptic principles.

Second Synthetical Reaction.—Mix together 1 part of boracic acid, 4 of acid tartrate of potassium, and 10 or 20 of water; evaporate to a syrupy consistence, spread on plates and set aside for dry scales to form. The resulting substance is, in water, far more readily soluble than either of its constituents, and is known as *boro-tartrate of potassium*, or *soluble cream of*

tartar. The Prussian *tartarus borazatus* differs from the foregoing French variety in containing 1 part of *borax* to 3 of acid tartrate of potassium.

Analytical Reactions (Tests).

First Analytical Reaction.—Dip a piece of turmeric paper (paper soaked in tincture of turmeric tubers and dried) into a solution of boracic acid; it is coloured brown-red, as by alkalies.

The usual way of applying this test is as follows:—Add to a solution of any borate a few drops of hydrochloric acid, immerse half of a slip of turmeric paper in the liquid, then remove the hydrochloric acid by drying the paper over a flame. Concentrated hydrochloric acid and ferric chloride produce a somewhat similar effect.

Second Analytical Reaction.—To a fragment of a borate or metaborate (*borax*, for example) in a small dish or watch-glass add a drop of sulphuric acid and then a little alcohol, warm the mixture and set light to the spirit; the resulting flame will be tinged of a greenish colour at its edges by the volatilized metaboracic acid or boracic anhydride.

The liquid should be well stirred while burning. Salts of copper and some metallic chlorides produce a somewhat similar colour. The flame test may also be applied to a little of a mixture of the borate with strong sulphuric acid on a platinum wire. Glycerine may be used in place of sulphuric acid (115).

Other Analytical Reactions.—In solutions of *borax* barium salts give a white precipitate of barium metaborate ($\text{Ba}_2\text{B}_2\text{O}_5$) soluble in acids and alkaline salts. Nitrate of silver gives metaborate of silver (AgBO_2), soluble in nitric acid and in ammonia. Chloride of calcium, if the solution is not too dilute, gives white borate of calcium.

QUESTIONS AND EXERCISES.

571. Illustrate the relation of vanadium to nitrogen by formulæ of compounds of each element.

572. Describe the preparation of borax.

573. Give the formulæ of boracic acid, metaboracic acid, and borax.

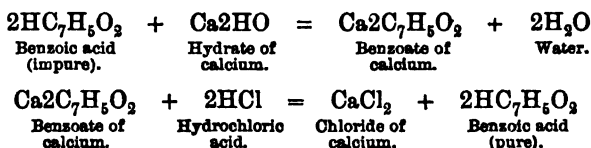
574. Mention the tests for borates or metaborates.

The foregoing acids and other salts contain the only acidulous radicals that are commonly met with in analysis or in ordinary medical or pharmaceutical operations. There are, however, many others which occasionally present themselves. The chief of these will now be shortly noticed; they are arranged in alphabetical order to facilitate reference.

SALTS OF RARER ACIDULOUS RADICALS.

BENZOIC ACID ($\text{HC}_7\text{H}_5\text{O}_2$) AND OTHER BENZOATES.—Slowly heat a fragment of benzoin (Gum Benjamin) (*Benzoinum*, B. P.) in a test-tube; benzoic acid (*Acidum Benzoicum*, B. P.) rises in vapour and condenses in small, white, feathery plates and needles on the cool sides of the tube. If the benzoin is first mixed with twice its weight of sand or roughly powdered pumice-stone, and the heat very cautiously applied, the product will be less likely to be burnt, and a larger quantity be yielded. By repeated sublimation 10 to 15 per cent. may be obtained.

A more economical process is to boil the benzoin with one-fourth its weight of lime, filter, concentrate, decompose the solution of benzoate of calcium by hydrochloric acid, collect the precipitated benzoic acid, press between paper, dry, and sublime in a tube or other vessel.



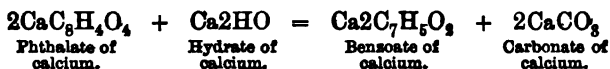
There is always associated with the product a minute

quantity of a volatile oil (styrol ?) of agreeable odour, suggesting that of hay.

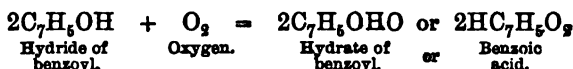
Benzoic acid is also prepared on a large scale artificially from naphthalin, one of the crystalline by-products in the distillation of coal for gas. The naphthalin is oxidized by nitric acid to naphthalic or phthalic acid :—



The phthalic acid is neutralized by lime and the phthalate of calcium heated with hydrate of calcium in a covered vessel at a temperature of about 640° F. for several hours. Benzoate and carbonate of calcium are formed, and, from the powder, the benzoic acid is set free by action of hydrochloric acid.

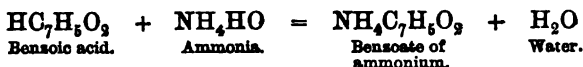


The crystalline deposit formed when essential oil of almonds (hydride of benzoyl or benzoic aldehyd) is exposed to the air is benzoic acid.



Benzoic acid is also obtained from hippuric acid (p. 396).

Benzoate of Ammonium.—To a little benzoic acid add a few drops of solution of ammonia; it readily dissolves, forming benzoate of ammonium (*Ammonia Benzoas*, B. P.) ($\text{NH}_4\text{C}_7\text{H}_5\text{O}_2$).



On evaporation, acid crystals or, ammonia being added, neutral crystals of benzoate of ammonium are deposited.

Properties.—Benzoic acid is also soluble in other alkaline liquids, forming benzoates. It is slightly soluble in cold water, more so in hot, and readily soluble in rectified spirit. It melts at 248° F. and boils at 462°, volatilizing with only a slight residue.

Tests.—The following are the *tests* for benzoic acid :—To

a portion of the above solution of benzoate of ammonium add a drop or two of sulphuric or hydrochloric acid; a white crystalline precipitate of benzoic acid separates. To another portion, carefully made neutral, add a drop or two of neutral solution of perchloride of iron; reddish ferric benzoate is precipitated.

Cinnamic acid ($C_9H_9O_2$).—Benzoic acid is distinguished from an allied body, cinnamic acid (occurring in Balsams of Peru, Tolu, and Storax, and sometimes in Benzoin), by not yielding hydride of benzoyl (C_7H_5OH) (oil of bitter almonds) when distilled with chromic acid—that is, with a mixture of red chromate of potassium and sulphuric acid.

CARMINE ACID ($C_{14}H_{14}O_8$).—This is the colouring principle of cochineal (*Coccus*, B. P.). The carmine of trade, when unadulterated (*vide Pharmaceutical Journal*, 1859–60, p. 546), is carminic acid united with one or two per cent. of alumina, or, occasionally, of oxide of tin or albumen. It should be wholly soluble in solution of ammonia, giving an apparently clear, rich, purple liquid. Carmine, with French chalk, or starch, constitutes *face-rouge* or *animal rouge*.

Merrick tests the relative value of several samples of cochineal or carmine by observing how much solution of permanganate of potassium is required to change the colour of a decoction to faint pink.

CETRARIAC ACID ($H_2C_{34}H_{30}O_{16}$) is the bitter principle of Iceland "moss" (*Cetraria*, B. P.). In the lichen it is associated with much starch.

CHRY SOPHANIC ACID ($C_{18}H_{10}O_4$).—This acid is the chief colouring-matter of various species of rhubarb-root (*Rhei Radix*, B. P.) and, under the name of *parietinic acid*, of various common yellow lichens. The author also finds it to form four-fifths of "Chrysarobine," a name given by Kemp to the pith, etc., of a leguminous tree. Chrysarobine is also known as Araroba Powder, Bahia Powder, Brazil Powder, Goa Powder, and Ringworm Powder. The chrysarobine as it occurs in the tree, or when fresh, has since been shown by Liebermann and Seidler to have the formula $C_{30}H_{26}O_7$; this, by oxidation and elimination of water, yields the chrysophanic acid. Chrysophanic acid may be obtained in crystals of a golden-yellow colour, hence the name (from *χρυσός*, *chrysos*,

gold, and *φαίρω*, *phainō*, I shine). Its synonyms are *Rhaponticin*, *Rheic acid*, *Rhein*, *Rheumin*, *Rhubarbaric acid*, *Rhubarbarin*, *Rumicin*. Chrysophanic acid, black, red-brown, and red resins (*Aporetine*, *Phæoretine*, and *Erythroretine*), a bitter principle, and tannic acid are considered to be the conjoint source of the therapeutic properties of rhubarb. Chrysophanic acid is also contained in several species of *Rumex* or *Dock*. "Rumicin" is a preparation of Yellow Dock.

CYANIC ACID (HCyO) AND OTHER CYANATES.—The valuable reducing-power of cyanide of potassium (KCy) (or ferrocyanide, K_4Fcy) on metallic compounds is due to the avidity with which cyanate (KCyO) is formed.

Process.—Fuse a few grains of cyanide of potassium in a small porcelain crucible, and add powdered oxide of lead; a globule of metallic lead is at once set free, excess of the oxide converting the whole of the cyanide of potassium into cyanate of potassium.

Urea.—Cyanate of potassium (KCNO), or, better, cyanate of lead (Pb_2CNO), treated with sulphate of ammonium, yields cyanate of ammonium (NH_4CNO); and solution of cyanate of ammonium, when simply heated, changes to artificial *urea* ($\text{CH}_4\text{N}_2\text{O}$), the most important constituent of urine, and the chief form in which the nitrogen of food is eliminated from the animal system. The process will be more fully described subsequently in connection with *urea*.

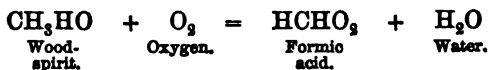
FORMIC ACID (HCHO_2).—The red ant (*Formica rufa*) and several other insects, when irritated, eject a strongly acid, acrid liquid, having a composition expressed by the above formula, and which has appropriately received the name of formic acid; it is also contained in the leaves of the stinging-nettle. (According to Church the sting of the *wasp* is *alkaline*.)

Process.—It may be artificially prepared by heating equal weights of oxalic acid and glycerine to a temperature of from 212°F . to 220° for fifteen hours. The glycerine has, apparently, no chemical action, but, for some unknown reason, induces decomposition of the oxalic acid at a lower tempera-

ture than would otherwise be necessary; at a higher temperature the formic acid itself is decomposed. On distilling the mixture with water the formic acid slowly passes over. The dilute acid may be concentrated by neutralizing with carbonate of lead, filtering, evaporating to a small bulk, collecting the deposited crystalline formate of lead, drying, decomposing in a current of sulphuretted hydrogen, separating the resulting syrupy acid, and passing dry air through the product until all sulphuretted hydrogen is removed.



Formic Acid may be instructively though not economically prepared by the oxidation of methylic alcohol (wood-spirit) just as acetic acid and valerianic acid are obtained from ethylic alcohol and amylic alcohol respectively.



Tests.—Formic acid does not char when heated alone or with sulphuric acid, but splits up into carbonic oxide gas and water. It is recognised by this property and by its reducing-action on salts of gold, platinum, mercury, and silver. It is solid below 32° F.

GALLIC ACID.—See *Tannic Acid*.

HEMIDESMIC ACID.—The supposed active principle of hemidesmus root (*Hemidesmi Radix*, B. P.).

HIPPURIC ACID ($\text{HC}_9\text{H}_8\text{NO}_3$) is a constituent of human urine (much increased on taking benzoic acid), but is best prepared from the urine of the horse (hence the name, from *ἵππος*, *hippos*, a horse), or, better, from that of the cow. To such urine add a little milk of lime, boil for a few minutes, remove precipitated phosphates by filtration, drop in hydrochloric acid until the liquid, after well stirring, is exactly neutral to test-paper, concentrate to about one-eighth the original bulk, and add excess of strong hydrochloric acid; impure hippuric acid is deposited. From a solution of the impure acid in hot water chlorine gas removes the colour,

and the liquid deposits crystals of pure hippuric acid on cooling. Its constitution is that of *benzoic glycocine*, $C_2H_2H_2(C_7H_5O)NO_2$.

Tests.—To a solution of hippurate add neutral solution of ferric chloride; a brown precipitate (ferric hippurate) results. Salts of silver and mercury give white precipitates. Heat hippuric acid in a test-tube; it chars, benzoic acid sublimes, and vapours of characteristic odour are evolved; they contain, amongst other bodies, hydrocyanic acid and a substance smelling somewhat like Tonka bean.—The crystalline form of hippuric acid is characteristic; it will be described in connection with the subject of urine.

QUESTIONS AND EXERCISES.

575. Give the preparation, composition, properties, and tests of benzoic acid, employing equations or diagrams.

576. What is the nature of carmine?

577. Name the bitter principle of Iceland "moss."

578. Mention the colouring principle of rhubarb.

579. To what is rhubarb considered to owe its medicinal activity?

580. How is cyanate of potassium prepared, how converted into an ammonium salt, and what are the relations of the latter to urea?

581. Give the formulæ of cyanic acid, cyanite of ammonium, and urea.

582. What is the chemical formula of formic acid?

583. Describe the artificial production of formic acid.

584. Describe the relation of formic acid to wood-spirit.

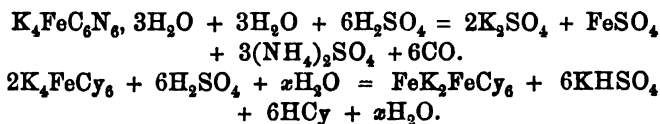
585. State the sources, characters, and tests of hippuric acid.

HYDROFERROCYANIC ACID ($H_4Fe^{\prime}Cy_6$, or H_4Fcy''') AND OTHER FERROCYANIDES.—The ferrocyanide of most interest is that of potassium, the "yellow prussiate of potash" (*Potassæ Prussias Flava*, B. P.) ($K_4FeC_6N_6 \cdot 3H_2O$, the formation of which was alluded to in connection with hydrocyanic acid (see p. 323). It cannot be regarded as simply a double salt of cyanide of potassium with cyanide of iron ($FeCy_3$, $4KCy$), its chemical properties being entirely different from either of those sub-

stances; moreover, unlike cyanide of potassium, it is not poisonous. Most of its reactions point to the conclusion that its iron and cyanogen are intimately united to form a definite quadrivalent radical appropriately termed *ferrocyanogen* (FeCy_6 , or Fcy). One part of ferrocyanide of potassium in 20 of water forms the official "Solution of Yellow Prussiate of Potash," B. P.

Tests.—Many of the ferrocyanides are insoluble, and are therefore precipitated when solution of ferrocyanide of potassium is added to the various salts. Those of iron and copper being of characteristic colour, are adopted as tests of the presence of the metals or of the ferrocyanogen, as the case may be. To solution of ferrocyanide of potassium add a ferric salt; ferrocyanide of iron (Fe_4Fcy_3) (Prussian blue) is precipitated. To another portion add solution of a copper salt; reddish-brown ferrocyanide of copper (Cu_2Fcy) is precipitated.

Note.—The ferrocyanogen in ferrocyanide of potassium is broken up when the salt is heated with sulphuric acid, carbonic oxide being evolved if the acid is strong (that is, ordinary oil of vitriol— H_2SO_4 with 3 to 4 per cent. of water), and hydrocyanic acid if weak:—



Hydrocyanic Acid has already been described. (*Vide* p. 324.)

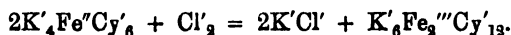
Carbonic Oxide (CO).—Heat two or three fragments of ferrocyanide of potassium with eight or ten times their weight of sulphuric acid, and, as soon as the gas begins to be evolved, remove the test-tube from the flame; for the action, when once set up, proceeds somewhat tumultuously. Ignite the carbonic oxide at the mouth of the tube; it burns with a pale blue flame, the product of combustion being carbonic acid gas (CO_2).

Carbonic oxide is a direct poison. It is generated whenever coke, charcoal, or coal burns with an insufficient supply

of air. Hence the danger of open fires in the more or less closed apartments of ordinary dwellings.

Carbonic oxide may also be obtained from oxalic acid. (*Vide* p. 370.)

HYDROFERRIDCYANIC ACID ($H_6Fe''_2Cy_{13}$, or $H^i_6Fdcy^{VI}$) AND OTHER FERRIDCYANIDES.—Pass chlorine gas slowly through solution of ferrocyanide of potassium until the liquid, after frequent shaking, ceases to give a blue precipitate when a minute portion is taken out on the end of a glass rod and brought into contact with a drop of a dilute solution of a ferric salt; it now contains ferridcyanide of potassium ($K_6Fe''_2Cy_{13}$, or $K^i_6Fdcy^{VI}$), *red prussiate of potash* (B. P.), as it is termed, from the colour of its crystals. Excess of chlorine must be carefully avoided, as chloride of cyangen and other compounds are then formed.



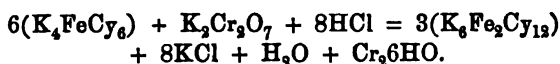
Another process.—To a cold solution of yellow prussiate of potash so much hydrochloric acid is added as will take two atoms of potassium from two molecules of the salt, and then a cold clear solution of bleaching-powder till ferric chloride gives no reaction. Any excess of acid is then neutralized with chalk, and the solution evaporated to crystallization (Rhien).

Note.—The removal of two atoms of potassium from the ferrocyanide molecules is the only change of composition that occurs; but the ferrocyanogen is altered in quality, its iron passing from the ferrous to the ferric condition, from bivalent to trivalent activity—altered to a condition in which it no longer precipitates ferric salts, but gives a dark-blue precipitate with ferrous salts.

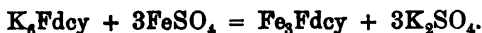
The radical is distinguished as *ferridcyanogen*.

Ferridcyanide of potassium may also be prepared by a modification of the foregoing method in which nascent instead of free chlorine is employed (Wenzell). Take of Bichromate of Potash 1 part, Ferrocyanide of Potassium Cryst. 5.72 parts, Hydrochloric Acid, of spec. grav. 1.16, 3 parts, by weight, Water 60 parts. Dissolve the two salts in hot water, add the acid, heat to boiling, continuing the ebullition, replacing the water evaporated during the process until a portion of the

filtered liquid is not precipitated on the addition of solution of ferric chloride. When reaction is completed, filter the liquid, and wash the hydrate of chromium, unite the liquids, and concentrate to crystallization. If the evaporated liquid possess an acid reaction, the addition of caustic potash, in sufficient quantity to cause a weak alkaline reaction, will greatly facilitate the subsequent crystallization.



Test.—To a portion of the solution add solution of ferrous sulphate; a precipitate falls. This precipitate is ferridcyanide of iron (Turnbull's blue), $Fe'_3Fe''_2Cy'_{12}$, or Fe''_3Fdcy^V .



It will be noticed that the change in the condition of the iron keeps up the balance of the atomic values of the various parts of the radicals or of the salts; the quantivalential equilibrium is maintained.

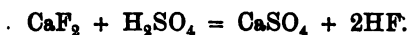
A solution of 1 of ferridcyanide of potassium in 20 of water constitutes the "Solution of Red Prussiate of Potash," B. P.

HYDROFLUORIC ACID (HF) AND OTHER FLUORIDES.—Molecular weight of HF, 20. The chief use of hydrofluoric acid is in etching on glass. The operation, performed on the small scale, also constitutes the best test for fluorine, the elementary radical of all fluorides.

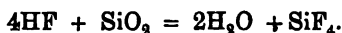
Process and Test.—Warm any odd piece of window-glass, having an inch or two of surface, until a piece of beeswax rubbed on one side yields a thin oily film. When cool make a cross, letter, or other mark on the glass by pressing a pointed piece of wood, a penknife, or file through the wax. Place a few grains of powdered fluor spar, the commonest natural fluoride, in a porcelain crucible (or a lead cup), add a drop or two of sulphuric acid, cover the crucible with the prepared glass, waxed side downwards, and gently warm the bottom of the crucible in a fume chamber or in the open air, in such a way as not to melt the wax. After a few minutes remove the glass, wash the waxed side by pouring water over

it, scrape off most of the wax, then warm the glass and wipe off the remainder; the marks made through the wax will be found to be permanently etched on the glass; the acid has eaten into or *etched* (from the German *ätzen*, to corrode) the glass.

In the above operation the fluoride of calcium and sulphuric acid yield hydrofluoric acid, thus:—



The hydrofluoric acid gas and the silica of the glass then yield gaseous fluoride of silicon (SiF_4), which escapes, and water thus:—



The silica being removed from the glass, leaves furrows or etched portions.

Note.—In the experiment just described, the liberated hydrofluoric acid also attacks the siliceous glazing of the porcelain crucible; so that in important cases, where search is made for very small quantities of fluorine, vessels of platinum or lead must be employed.

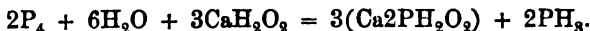
Uses.—The aqueous solution of hydrofluoric acid used by etchers, and commonly termed simply hydrofluoric acid, or “fluoric” acid, is prepared in leaden stills and receivers, and kept in leaden or gutta-percha bottles. Except these materials, as well as platinum and fluor spar, hydrofluoric acid rapidly attacks any substance of which bottles and basins are usually made. It quickly cauterizes the skin, producing a painful slow-healing sore.

Quantivalence.—The atom of fluorine, like that of chlorine, bromine, or iodine, is univalent ($\text{F}^{\cdot}$). The great analogy existing between these radicals extends to their compounds.

Fluorine is said to be a colourless gas; but, from the avidity with which it combines with all elements (except oxygen), it is so difficult of isolation as hitherto to preclude satisfactory study of its physical properties.

HYPOPHOSPHOROUS ACID (H_3PO_2 , or HPH_2O_2) AND OTHER HYPOPHOSPHITES.—Boil together, in a fume chamber, a grain or two of phosphorus, a few grains of slaked lime, and about a quarter of an ounce of water, until phosphoretted hydrogen, a spontaneously inflammable, badly smelling gas, ceases to be

evolved. The mixture, filtered, and excess of lime removed by carbonic acid gas, yields solution of hypophosphite of calcium ($\text{Ca}_2\text{PH}_2\text{O}_3$) (*Calci Hypophosphis*, B. P.).

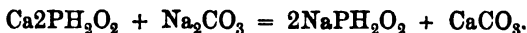


The solution, when concentrated by evaporation, has been known to explode, probably from formation of phosphoretted hydrogen. This may be prevented, it is said, by evaporating at a low temperature, especially towards the close of the operation; or by adding alcohol, which decomposes any traces of liquid phosphoretted hydrogen (PH_3) or solid phosphoretted hydrogen (P_2H) which possibly may be present, and to which it is conceivable explosion may be due.

Phosphoretted hydrogen (PH_3).—The above reaction is also that by which phosphoretted hydrogen, the third hydride of phosphorus, may be prepared. If the gas is to be collected, the phosphorus, and water, may first be boiled in a flask until a jet of spontaneously inflammable phosphorus vapour escapes, with steam, from the end of the attached delivery-tube. Strong hot solution of caustic potash or soda is next very gradually poured into the flask through a funnel tube previously fitted into the cork, the liquid being kept boiling. Phosphoretted hydrogen is then evolved, and if the delivery-tube dip under water may be collected, or allowed to slowly pass up through the water bubble by bubble so as to form the peculiar rings of smoke (phosphoric anhydride) characteristic of the experiment.

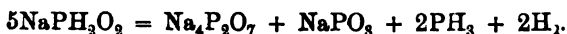
Hypophosphite of calcium (*Calci Hypophosphis*, U. S. P.) may be obtained in crystals by evaporation and slow cooling; but the solution is usually at once evaporated to dryness, a white pulverulent salt being obtained. Other hypophosphites may be obtained in the same way from other hydrates, or by double decomposition of the calcium salt and carbonates.

Hypophosphite of sodium (NaPH_2O_3 , H_2O) (*Sodæ Hypophosphis*, B. P.) is made by decomposing solution of hypophosphite of calcium by carbonate of sodium, filtering, and evaporating to dryness. It is a white, granular, deliquescent substance.



When heated, the water is first evolved, then hydrogen and spontaneously inflammable phosphoretted hydrogen, and a

mixture of pyrophosphate and metaphosphate of sodium remains (Rammelsberg).



Hypophosphorous acid, the hydrogen hypophosphite, may be prepared by decomposing the calcium salt by oxalic acid; hypophosphite of quinine by dissolving the alkaloid in hypophosphorous acid, or by decomposing sulphate of quinine by hypophosphite of barium.

The hypophosphites are often used in medicine in the form of syrups. The term hypophosphite is in allusion to the smaller amount (*ὑπό, hupo*, under or deficiency) of oxygen in these compounds (R_3PO_2) than in the phosphites (R_3PO_3), a class of salts having again less oxygen in their molecules than exists in those of the phosphates (R_3PO_4). The prefix *hypo* has similar significance in such words as hyposulphite and hypochlorite.

Tests.—To a portion of the above solution of hypophosphite of calcium add solution of chloride of barium, chloride of calcium, or acetate of lead; in neither case is a precipitate obtained, whereas soluble phosphates and phosphites yield white precipitates of phosphate or phosphite of barium, calcium, or lead. To other portions add solutions of nitrate of silver and mercuric chloride; the respective metals are precipitated as by phosphites. To another small portion add zinc and dilute sulphuric acid; hydrogen and phosphoretted hydrogen are evolved as from phosphites. To another portion add sufficient oxalic acid to remove the calcium; filter; to the solution of hypophosphorous acid add solution of sulphate of copper and slowly warm the mixture; solid brown cuprous hydride (Cu_2H_2) is precipitated: increase the heat to the boiling-point; hydrogen is evolved and metallic copper set free. Heat a small quantity of a solid hypophosphite on the end of a spatula in a flame; it splits up into pyrophosphate, a little metaphosphate, hydrogen, phosphoretted hydrogen, and, sometimes, water, burning with a phosphorescent light, the official hypophosphite of calcium yielding about 80 per cent. of residue.



To an acid solution of a hypophosphite add solution of a permanganate. It is instantly decolorized. The same effect follows its addition to an acid solution of a phosphite, but not to that of an ortho-, meta-, or pyrophosphate.

HYPOSULPHUROUS ACID ($\text{H}_2\text{S}_2\text{O}_3$) AND OTHER HYPOSULPHITES.

—The only hyposulphite of much interest in pharmacy is the sodium salt (Hyposulphite of Soda, B. P.) ($\text{Na}_2\text{S}_2\text{O}_3, 5\text{H}_2\text{O}$). Hyposulphites may be termed *Thiosulphates* (e. g. $\text{H}_2\text{SO}_3\text{S}$).

Process.—Heat together gently, or set aside in a warm place, a mixture of solution of sulphite of sodium (Na_2SO_3) and a little powdered sulphur; combination slowly takes place, and hyposulphite of sodium is formed. The solution, filtered from excess of sulphur, readily yields crystals. (The solution of sulphite of sodium may be made by saturating solution of soda with sulphurous acid gas.)

Use of hyposulphite of sodium in quantitative analysis.—In the British Pharmacopœia hyposulphite of sodium is given as a reagent for the quantitative estimation of free iodine in volumetric analysis. To a few drops of iodine-water add cold mucilage of starch; a deep-blue colour (starch iodide) is produced. To the product add solution of hyposulphite of sodium until the blue colour just disappears. This absorption of iodine is sufficiently definite and delicate to admit of application for quantitative purposes. It depends on the combination of the iodine with half of the sodium in two molecules of the hyposulphite, the hyposulphurous radicals of the two molecules apparently coalescing to form a new radical, the tetrathionic (from *τέρας*, *tēras*, four, and *θειον*, *theion*, sulphur),—tetrathionate of sodium ($\text{Na}_2\text{S}_4\text{O}_6$) and iodide of sodium being formed.

Sulphur oxyacids.—It will be as well here to give the formulæ of other oxyacids of sulphur, forming with the four already mentioned a series that is as useful as the series of compounds of nitrogen and oxygen in illustrating the soundness of Dalton's atomic theory.

Hydrosulphurous Acid H_2SO_2	Dithionic Acid . . $\text{H}_2\text{S}_2\text{O}_6$
Sulphurous Acid . . H_2SO_3	Trithionic Acid . . $\text{H}_2\text{S}_3\text{O}_6$
Sulphuric Acid . . H_2SO_4	Tetrathionic Acid . $\text{H}_2\text{S}_4\text{O}_6$
Hyposulphurous Acid $\text{H}_2\text{S}_2\text{O}_3$	Pentathionic Acid . $\text{H}_2\text{S}_5\text{O}_6$

Use of "Hypo" in Photography.—The sodium hyposulphite is largely used in photography to dissolve chloride, bromide, or iodide of silver off plates which have been exposed in the camera. Prepare a little chloride of silver by adding a chloride (chloride of sodium) to a few drops of solution of nitrate of silver. Collect the precipitated chloride on a filter, wash, and add a few drops of solution of hyposulphite of sodium; the silver salt is dissolved, solution of double hyposulphite of sodium and silver being formed. The solution of this double hyposulphite has a remarkably sweet taste, sweeter than syrup if the solution is strong. The double hyposulphite of sodium and gold is employed for giving a pleasant tint to photographic prints.

Test.—To solution of a hyposulphite add a few drops of dilute sulphuric or other acid; hyposulphurous acid is set free, but at once begins to decompose into sulphurous acid, recognised by its odour, and free sulphur ($2\text{H}_2\text{S}_2\text{O}_3 = 2\text{H}_2\text{SO}_3 + \text{S}_2$). This reaction constitutes the best test for hyposulphites. Another good test of a soluble simple hyposulphite is its power of dissolving chloride of silver with production of a more or less sweet solution.

QUESTIONS AND EXERCISES.

586. Give the formula of ferrocyanide of potassium.
587. What is the supposed constitution of ferrocyanide of potassium?
588. Enumerate the tests for ferrocyanogen.
589. What are the respective reactions of ferrocyanide of potassium with strong and weak sulphuric acid?
590. Mention and explain a common source of carbonic oxide in households. What is the product of its combustion?
591. Write equations or diagrams illustrative of the changes

effected on ferrocyanide of potassium during its conversion into ferridcyanide.

592. By what reactions may the presence of a ferridcyanide in a solution be demonstrated?

593. State the difference between Prussian blue and Turnbull's blue.

594. Describe the source, mode of preparation, chief use of, and test for hydrofluoric acid.

595. Illustrate by a diagram the preparation and composition of hyposulphite of sodium.

596. Mention the uses and characteristic reactions of hyposulphite of sodium.

597. Give the names and formulæ of eight acids, each containing hydrogen, sulphur, and oxygen.

LACTIC ACID ($\text{H}_3\text{C}_3\text{H}_4\text{O}_3$) AND OTHER LACTATES.—Lactic acid occurs naturally in willow bark (Dott). When milk turns sour its sugar has become converted into an acid appropriately termed lactic (*lac*, *lactis*). Other saccharine and amylaceous substances also by fermentation yield lactic acid. Neither the hydrogen lactate (lactic acid) nor other lactates are much used in England.

Process.—Lactate of calcium and lactic acid may be prepared as follows:—Mix together eight parts of sugar, one of common cheese, three of chalk, and fifty of water, and set aside in a warm place (about 80°F .) for two or three weeks; a mass of small crystals of lactate of calcium results. Remove these, recrystallize from hot water, decompose by sulphuric acid, avoiding excess, digest in alcohol, filter off the sulphate of calcium, evaporate the clear solution to a syrup; this residue is lactic acid. Sp. gr. 1.212.

Test.—No single reaction of lactic acid is sufficiently distinctive to form a test. The crystalline form of the lactate of calcium, as seen by the microscope, is characteristic. The production of this salt, and the isolation of the syrupy acid itself, are the only means, short of quantitative analysis, on which reliance can be placed.

A variety of lactic acid has been obtained from the juice of

flesh ; it is termed *sarcolactic acid* (from $\sigma\alpha\rho\acute{\epsilon}\varsigma$, $\sigma\alpha\rho\kappa\acute{o}\varsigma$, *sarr*, *sarcos*, flesh).

MALIC ACID ($\text{H}_3\text{C}_4\text{H}_3\text{O}_5$) AND OTHER MALATES (from *malum*, an apple).—The juice of unripe apples, gooseberries, currants, rhubarb-stalks, etc., contains malic acid and malate of potassium. When isolated it occurs in deliquescent prismatic crystals.

Tests.—Malate of calcium ($\text{CaHC}_4\text{H}_3\text{O}_5$) is soluble in water ; hence the aqueous solution of malic acid or other malate is not precipitated by lime-water or chloride of calcium ; but on adding spirit of wine a white precipitate falls, owing to the insolubility of the calcium malate in alcohol. Malates are precipitated by lead-salts ; on warming the malate of lead with acetic acid it dissolves, separating out in acicular crystals on cooling. If the mixture be heated without acid the malate of lead agglutinates and fuses.

Hot strong sulphuric acid chars malic acid far less readily than it does nearly all other organic acids.

Asparagin ($\text{C}_4\text{H}_8\text{N}_2\text{O}_3, \text{H}_2\text{O}$).—This proximate principle of plants occurs in many vegetable juices, and doubtless plays a very important part in their nutrition. It is deposited in crystals when the fresh juices of asparagus, marsh-mallow, etc., are rapidly evaporated. It is noticed here because malic acid is readily obtained from it by oxidation, nitrogen being eliminated, and because its exact natural position among chemical substances is not yet well made out. The atoms of its molecule are those of aspartate of ammonium ($\text{NH}_4\text{C}_4\text{H}_5\text{NO}_4$), into which it is converted when its solution is long boiled. Decomposed by aid of ferments, asparagin, absorbing hydrogen, yields succinate of ammonium (NH_4)₂ $\text{C}_4\text{H}_4\text{O}_4$. Such reactions as these and the formation of lactic acid from sugars should suggest to the student possible modes in which chemical changes take place in the plant-department of the vast laboratory of nature.

MECONIC ACID ($\text{H}_3\text{C}_7\text{HO}_7$).—Opium contains meconic acid (from $\mu\acute{\eta}\kappa\omega\nu$, $\mu\acute{\epsilon}\lambda\kappa\acute{o}\nu$, a poppy) partially combined with morphia. To concentrated infusion of opium, nearly neutralized with ammonia, add solution of chloride of calcium ; meconate

of calcium is precipitated. Wash the precipitate, place it in a small quantity of hot water, and add a little hydrochloric acid; the clear liquid (filtered, if necessary) deposits scales of meconic acid on cooling.

Tests.—To solution of meconic acid or other meconate, or to infusion of opium, add a *neutral* solution of ferric chloride; a red solution of meconate of iron is produced. To a portion of the mixture add solution of corrosive sublimate; the colour is not destroyed: to another portion add hydrochloric acid; the colour is discharged. (These reagents act on sulphocyanate of iron, which is of similar tint, in exactly the opposite manner.)

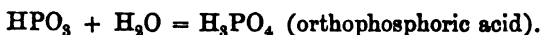
The normal meconates of potassium, sodium, and ammonium are soluble in water, the acid meconates very slightly soluble, the meconates of barium, calcium, lead, copper, and silver insoluble in water, but soluble in acetic acid.

METAPHOSPHORIC ACID (HPO_3) AND OTHER METAPHOSPHATES.—Prepare phosphoric anhydride (P_2O_5) by burning a small piece of phosphorus in a porcelain crucible placed on a plate and covered by an inverted test-glass, tumbler, half-pint measure-glass, or some such vessel. After waiting a few minutes for the phosphoric anhydride to fall, pour a little water on the plate and filter the liquid; the product is solution of metaphosphoric acid (from *μετά, meta*, a preposition denoting change).

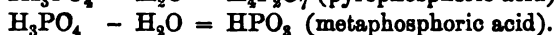


Tests.—To solution of metaphosphoric acid add ammonio-nitrate of silver, or to a neutral metaphosphate add solution of nitrate of silver; a white precipitate (AgPO_3) is obtained. This reaction sufficiently distinguishes metaphosphates from the ordinary phosphates or orthophosphates (from *ὀρθός, orthos*, straight), as the common phosphates may, for distinction, be termed (which give, it will be remembered, a *yellow* precipitate with nitrate of silver). Another variety of phosphates shortly to be considered, the pyrophosphates, also give a white precipitate with nitrate of silver. To the solution of metaphosphoric acid obtained as above, or by the action of

acetic acid on a metaphosphate, add an aqueous solution of white of egg; coagulation of the albumen ensues. Neither orthophosphoric nor pyrophosphoric acid coagulates albumen. Boil the aqueous solution of metaphosphoric acid for some time; on testing the solution the acid will be found to have been converted into orthophosphoric acid:—



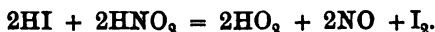
The ordinary medicinal phosphoric acid is made from phosphorus and nitric acid, the liquid being evaporated to a syrupy consistence to remove the last traces of nitric acid. It may contain pyrophosphoric and metaphosphoric acids, if the heat employed be high enough to remove the elements of water:—



On redilution the metaphosphoric acid only slowly reabsorbs water. If, therefore, on testing, metaphosphoric be found to be present, the solution should be boiled until conversion to orthophosphoric acid has occurred.

NITROUS ACID (HNO_2) AND OTHER NITRITES.—Strongly heat a fragment of nitrate of potassium or of sodium on a piece of platinum foil; oxygen is evolved and nitrite of potassium remains.

Test.—Dissolve the residue in water, add a few drops of dilute sulphuric acid, then a little weak solution of iodide of potassium, and, lastly, some mucilage of starch; the deep-blue compound of iodine and starch is at once produced. Repeat this experiment, using nitrate of potassium instead of nitrite; no blue colour is produced.



Test for Nitrites in Water.—This liberation of iodine by nitrites and not by nitrates is a reaction of considerable value in searching for nitrites in ordinary drinking waters, the occurrence of such salts being held to indicate the presence of nitrogenous organic matter in a state of oxidation or decay.

The sulphuric acid used in the operation must be pure, and the iodide of potassium free from iodate. If much organic matter is present, however, the nitric acid liberated by the sulphuric may be reduced to nitrous acid. It is perhaps best, therefore, to add acetic acid, and (Fresenius) distil over 10 or 20 per cent. of the water and apply the test to this distillate. Very dilute solutions of nitrous acid may thus be distilled without the slightest decomposition.

Commercial Nitrous Acid.—The liquid commonly termed in pharmacy "nitrous acid" is simply nitric acid impure from the presence of nitrous acid.

The only nitrite used in medicine is a nitrite of an organic basylous radical, ethyl; nitrite of ethyl ($C_2H_5NO_2$), or nitrous ether, is one of the chief constituents of "sweet spirit of nitre" (*Spiritus Ætheris Nitrosi*, B. P.: vide Index).

OPHELIC ACID ($C_{13}H_{20}O_{10}$).—This is one of the principles to which the herb *Ophelia chirata*, or Chiretta (*Chirata*, B. P.), owes its bitterness. It is an amorphous yellow body. Another is *Chiratin* ($C_{28}H_{48}O_{15}$), decomposable by hydrochloric acid into *Chiratogenin* ($C_{13}H_{24}O_8$) and ophelic acid (Höhn).

PHOSPHOROUS ACID (H_3PO_3 , or H_2PHO_3).—It is necessary to notice this compound in order that the reader may have brought before him the three acids of phosphorus, namely, phosphoric acid (H_3PO_4), phosphorous acid (H_2PHO_3), and hypophosphorous acid ($H_2PH_2O_3$): it will be noticed that in composition they differ from each other simply in the proportion of oxygen, the molecules containing four, three, and two atoms respectively. In constitution they differ by the hypothetical phosphoric radical or grouping being trivalent, the phosphorous radical bivalent, and the hypophosphorous radical univalent. These three acids and corresponding salts must not be confounded with pyrophosphoric and metaphosphoric acids and salts: the former are acids of phosphorus; the latter varieties of phosphoric acid: the former, in composition, differ from each other in the proportion of oxygen they contain; the latter by the elements of water:—

Acids of phosphorus.

H_3PO_4 phosphoric acid.

H_2PHO_3 phosphorous acid.

$H_2PH_2O_3$ hypophosphorous acid.

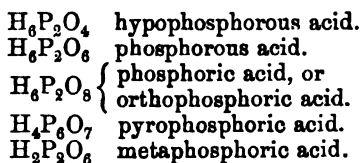
Varieties of phosphoric acid.

H_3PO_4 (ortho)phosphoric acid.

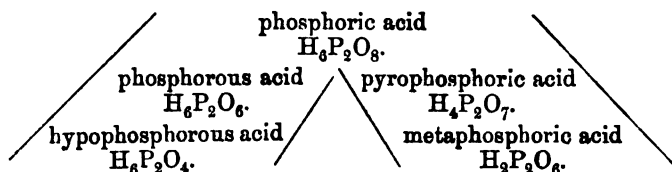
$H_4P_2O_7$ pyrophosphoric acid.

HPO_3 metaphosphoric acid.

When hypophosphorous acid is exposed to the air, oxygen is absorbed and phosphorous acid results; by prolonged exposure more oxygen is absorbed and phosphoric acid is obtained. When phosphoric acid, or rather, for distinction, orthophosphoric acid, is heated, every two molecules yield the elements of a molecule of water, and pyrophosphoric acid results; by prolonged exposure to heat more water is evolved, and metaphosphoric acid is obtained. These differences will be further evident if the formulæ be written empirically, nearly all being doubled, thus:—



Or thus:—



From the central compound, phosphoric acid, the acids of phosphorus differ by regularly diminishing proportions of the element oxygen (see previous page), the varieties of phosphoric acid by regularly diminishing proportions of the elements of water.

Prepare phosphorous acid by exposing a moist stick of phosphorus to the air; a thin stream of heavy white vapour falls, which contains the acid in question. The best method of collection is to place the stick in an old test-tube having a hole in the bottom, to support this tube by a funnel or otherwise, the neck of the funnel being supported in a bottle, test-glass, or tube, at the bottom of which is a little water. Having collected some phosphorous acid in this way, apply the various tests already alluded to under *Hypophosphorous Acid*, first carefully neutralizing the phosphorous acid by an alkali. The means by which the varieties of phosphoric acid are distin-

guished have been given under *Metaphosphoric Acid*. Associated with the phosphorous acid, prepared as above stated, there is said to be an acid having the formula H_2PO_3 and termed *hypophosphoric acid*. Its anhydride would be P_2O_4 .

Other soluble phosphites are prepared by neutralizing phosphorous acid with alkalis, and the insoluble phosphites by double decomposition.

PYROGALLIC ACID.—See Tannic Acid.

PYROPHOSPHORIC ACID ($H_4P_2O_7$) AND OTHER PYROPHOSPHATES.—Heat ordinary phosphate of sodium ($Na_2HPO_4, 12H_2O$) in a crucible; water of crystallization is first evolved, and dry phosphate (Na_2HPO_4) remains. Continue the heat to redness; two molecules of the salt yield one molecule of water, and a salt having new properties is obtained:—



It is termed pyrophosphate of sodium, in allusion to its origin (*πῦρ* *pῦr*, fire). Phosphoric acid itself is similarly affected by heat:— $2H_3PO_4 - H_2O = H_4P_2O_7$ (pyrophosphoric acid), though metaphosphoric acid is also formed. Other pyrophosphates are produced in a similar way, or by double decomposition and precipitation, or by neutralizing pyrophosphoric acid by an oxide, hydrate, or carbonate. Possibly the pyrophosphates are only compounds of orthophosphates with metaphosphates:—



Tests.—To solution of a pyrophosphate add solution of nitrate of silver; white pyrophosphate of silver ($Ag_4P_2O_7$) falls as a dense white powder, differing much in appearance from the white gelatinous metaphosphate of silver or the yellow orthophosphate. To pyrophosphoric acid, or to a pyrophosphate mixed with acetic acid, add an aqueous solution of albumen (white of egg); no precipitate occurs. Metaphosphoric acid, it will be remembered, gives a white precipitate with albumen.

QUESTIONS AND EXERCISES.

598. What are the sources of lactic acid ?
599. How is lactic acid usually prepared ?
600. Name some of the plants in which malic acid is found.
601. Whence is meconic acid derived ?
602. By what process may meconic acid be isolated ?
603. Which is the best test for the meconic radical ?
604. Distinguish meconates from sulphocyanates.
605. Give the mode of manufacture of hypophosphites.
606. How is phosphoretted hydrogen prepared ?
607. By what ready method may metaphosphoric acid be obtained for experimental purposes ?
608. Name the tests for metaphosphates.
609. How may meta- or pyro-phosphoric acid be converted into orthophosphoric acid ?
610. Describe the preparation of phosphorous acid.
611. State the relations which the acids of phosphorus bear to each other.
612. How are pyrophosphates prepared ?
613. Offer two views of the constitution of pyrophosphates.
614. Define, by formulæ, metaphosphates, pyrophosphates, orthophosphates, phosphites, and hypophosphites.
615. Mention the tests by which meta-, pyro-, and orthophosphates are analytically distinguished.
616. Name the reactions by which hypophosphites and phosphites are detected.

SILICIC ACID (H_4SiO_4) AND OTHER SILICATES.—Silicates of various kinds are among the commonest of minerals. The various *clays* are aluminium silicates ; the volcanic substance termed *pumice-stone* is a porous silicate of aluminium and of alkali-metals or of alkaline-earth metals ; *meerschaum* is an acid silicate of magnesium ; the ordinary *sandstones* are chiefly silica ; *sand*, *flint*, *quartz*, *agate*, *chalcedony*, and *opal* are silicic anhydride or *silica* (SiO_2). *Tripoli*, a polishing powder now found in many other countries than Tripoli, consists of infusorial skeletons of nearly pure silica. *Bath brick*, used in knife-polishing, is a silico-calcareous deposit found in the estuary at Bridgwater and other places. *Asbestos* or *amianth* is a fibrous silicate of calcium and magnesium, the length of the fibres being from less than one inch to five

feet. A single silk-like fibre can easily be fused, but, even in very small masses, and for all practical purposes, asbestos is infusible, and, of course, incombustible. It is also a bad conductor of heat. It is already largely used in packing piston rods and joints, and for steam apparatus generally; as a covering for boilers to prevent loss of heat by radiation; and for so lining ceilings, floors, and other partitions as to render rooms, etc., fireproof. Artificial silicates are familiar under the forms of *glass* and *earthenware*. Common English window-glass is usually silicate of calcium, sodium, and aluminium; French glass, silicate of calcium and sodium; Bohemian, chiefly silicate of potassium and calcium; English flint or crystal-glass for ornamental, table, and optical purposes, is mainly silicate of potassium and lead. Earthenware is mostly silicate of aluminium (clay), with more or less of silicate of calcium, sodium, and potassium, and in the commoner forms, silicate of iron. The various kinds of *porcelain* (China, Sèvres, Meissen, Berlin, English), *Wedgwood-ware*, and *stoneware* are varieties of earthenware. *Crucibles*, *bricks*, and *tiles* are clay-silicates. *Mortar* is essentially silicate of calcium. *Portland*, *Roman*, and other hydraulic cements* are silicates of calcium with more or less silicate of aluminium.

Mix together a few grains of powdered flint or sand with about five or six times its weight of carbonate of sodium and an equal quantity of carbonate of potassium, and fuse a little of the mixture on platinum-foil in the blowpipe-flame; the product is a kind of *soluble glass*. Boil the foil in water for a few minutes, filter; to a portion add excess of hydrochloric acid, evaporate the solution to dryness, and again boil the residue in water and acid; oxide of silicon, silicic anhydride, or *silica* (SiO_2) remains as a light, flaky, insoluble powder.

The *soluble glass* or *glass liquor* of trade commonly contains 10 or 12 per cent. of soda (NaHO) to 20 or 25 per cent. of silica (SiO_2).

The foregoing operation constitutes the *test* for silicates. By fusion with alkali the silicate is decomposed, and a soluble alkaline silicate formed. On addition of acid, silicic acid (H_4SiO_4) is set free, but remains in solution if sufficient

* For an article on "Cements" generally see the *Pharmaceutical Journal* for April 17th, 1880.

water is present. The heat subsequently applied eliminates water and reduces the silicic acid to silica (SiO_2), which is insoluble in water or acid. By the addition of hydrochloric acid to soluble glass, and removal of the resulting alkaline chloride and excess of hydrochloric acid by dialysis (a process to be subsequently described), a pure aqueous solution of silicic acid may be obtained; it readily changes into a gelatinous mass of silicic acid. Possibly some of the natural crystallized varieties of silica may have been obtained from the silica contained in such an aqueous solution, nearly all waters yielding a small quantity of silica when treated as above described.

A variety of silicic acid (H_2SiO_3), sometimes termed *dibasic*, to distinguish it from the normal or *tetrabasic* acid (H_4SiO_4), results when the aqueous solution of the latter is evaporated *in vacuo*.

Siliciuretted hydrogen, or hydride of silicon (SiH_4), is a spontaneously inflammable gas formed on treating silicide of magnesium with hydrochloric acid. It is the analogue of light carburetted hydrogen (CH_4). A liquid chloride of silicon (SiCl_4) and a gaseous fluoride (SiF_4) also exist.

Many other analogies are traceable between the elements silicon, boron, and carbon.

SUCCINIC ACID ($\text{H}_2\text{C}_4\text{H}_4\text{O}_4$).—Amber (*Succinum*) is a peculiar resin usually occurring in association with coal and lignite. From the fact that fragments of coniferous fruit are frequently found in amber, and impressions of bark on its surface, it is considered to have been an exudation from a species of *Pinus* now probably extinct. Heated in a retort, amber yields, first, a sour aqueous liquid containing acetic acid and another characteristic body appropriately termed *succinic acid*; second, a volatile liquid known as *oil of amber*, resembling the oil yielded by most resinous substances under similar circumstances; and, third, a pitchy residue allied to asphalt. The succinic acid is a normal constituent of the amber, the acetic acid is produced during distillation. Succinic acid has also been found in wormwood, in several pine-resins, and in certain animal fluids, such as those of hydatid cysts and hydrocele. It may be obtained artificially from butyric, stearic, or margaric acid by oxidation. Tartaric, malic, and succinic acids are also convertible the one into the other.

The *succinates* are normal ($R'_2C_4H_4O_4$) and acid ($R'HC_4H_4O_4$); a double succinate of potassium and hydrogen ($KHC_4H_4O_4$, $H_2C_4H_4O_4$, H_2O), analogous to the superacid oxalate, salt of sorrel, also exists.

Soluble succinates give a bulky brown precipitate with neutral ferric chloride, only less voluminous than ferric benzoate; a white precipitate with acetate of lead, soluble in excess of either reagent; with nitrate of silver a white precipitate after a time; with chloride of barium no precipitate at first, but a white one of succinate of barium on the addition of ammonia and alcohol. Succinates are distinguished from benzoates by the last-named reaction, and by not yielding a precipitate on the addition of acids (*vide* p. 394).

SULPHOCYANIC ACID (HCyS) AND OTHER SULPHOCYANATES.—Boil together sulphur and solution of cyanide of potassium; solution of sulphocyanate of potassium (KCyS) is formed. Warm the liquid, add hydrochloric acid till it faintly reddens litmus-paper, and filter; any sulphide of potassium is thus decomposed, and the solutions may then be used for the following reactions.

Tests.—To a small portion of the solution add a ferric salt (Fe_2Cl_6); a deep-blood-red solution of ferric sulphocyanate is formed. To a portion of the red liquid add a little hydrochloric acid; the colour is not discharged (meconate of iron, a salt of similar tint, is decomposed by hydrochloric acid). In the acid liquid place a fragment or two of zinc; sulphuretted hydrogen is evolved, and the red colour disappears. To another portion of the ferric sulphocyanate add solution of corrosive sublimate; the colour is at once discharged. (Ferric meconate is unaffected by corrosive sublimate.) The ferric is the best test of the presence of a sulphocyanate; indirectly, it is a good test of the presence of hydrocyanic acid or cyanogen. Solutions of pure ferrous salts are not coloured by the solution of sulphocyanate.

To solution of a sulphocyanate add solution of mercuric

nitrate; mercuric sulphocyanate is precipitated as a white powder.

Pharaoh's Serpents.—Mercuric sulphocyanate, thoroughly washed and made up into little cones, forms the toy termed Pharaoh's serpent. It readily burns when ignited, the chief product being a light solid matter (mellon, C_9N_{13} , and melam, $C_3H_6N_6$), which issues from the cone in a snake-like coil of extraordinary length. The other products are mercuric sulphide (of which part remains in the snake and part is volatilized), nitrogen, sulphurous and carbonic acid gases, and vapour of metallic mercury. (For details concerning the economical manufacture of sulphocyanates see *Pharmaceutical Journal*, 2nd series, vol. vii. p. 581, and p. 152.)

The sulphocyanic radical (CyS) is often termed sulphocyanogen (Soy), and its compounds regarded as *sulphocyanides*. Saliva contains sulphocyanates.

TANNIC ACID OR TANNIN ($C_{27}H_{22}O_{17}$, or perhaps $C_{11}H_{12}O_9$). This is a common astringent constituent of plants, but is contained in largest quantity in galls (excrecences on the oak formed by the puncture and deposited ova of an insect). English galls contain from 14 to 28 per cent. of tannic acid; Aleppo galls (*Galla*, B. P.) from 25 to 65 per cent. (*Acidum Tannicum*, B. P.).

Process.—"Expose powdered galls (about an ounce is sufficient for the purpose of study) to a damp atmosphere for two or three days, and afterwards add sufficient ether to form a soft paste. Let this stand in a well-closed vessel for twenty-four hours, then, having quickly enveloped it in a linen cloth, submit it to strong pressure so as to separate the liquid portion, which contains the bulk of the tannin in solution. Reduce the pressed cake to powder, mix it with sufficient ether, to which one-sixteenth of its bulk of water has been added, to form again a soft paste, and press this as before. Mix the expressed liquids, and expose the mixture to spontaneous evaporation until, by the aid subsequently of a little heat, it has acquired the consistence of a soft extract; then place it on earthen plates or dishes, and dry it in a hot-air chamber at a temperature not exceeding 212° F."

The resulting tannic acid occurs in "pale yellow vesicular masses or thin glistening scales, with a strongly astringent taste and an acid reaction, readily soluble in water and rectified spirit, very sparingly soluble in pure ether," though soluble in the ethereal fluid used in the foregoing process—a fluid which is really a mixture of true ether, water and alcohol (both the latter contained in the common "ether") and a little added water also.

Medicinal Uses.—Tannic acid is very soluble in water, and in this form is usually administered in medicine. Its official preparations are *Glycerinum Acidi Tannici*, *Suppositoria Acidi Tannici*, and *Trochisci Acidi Tannici*.

Tests.—To an aqueous solution of tannic acid add aqueous solution of gelatine; a yellowish-white flocculent compound of the two substances is precipitated. This is a good test of the presence of tannic acid.

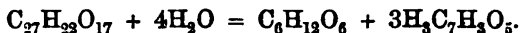
Tanning.—The above reaction also serves to explain the chemical principle involved in *tanning*—the operation of converting skin into leather. In that process the skin is soaked in infusion of oak-bark (*Quercus Cortex*, B. P.), the tannic acid of which uniting with the gelatinous tissues of the skin yields a compound very well represented by the above precipitate. The outer bark of the oak contains little or no tannic acid, and is commonly shaved off from the pieces of bark which are large enough to handle; useless colouring-matter is thus also rejected. Other infusions and extracts besides that of oak-bark (chiefly catechu, sumach, and valonia) are largely used by tanners; if used alone these act too quickly, and give a harsh, hard, less durable leather. The tannic acid of these preparations is probably slightly different from that of oak-bark.

To an aqueous solution of tannic acid add a neutral solution of a ferric salt; dark bluish-black tannate of iron is slowly precipitated. This is an excellent test for the presence of tannic acid in vegetable infusions. The precipitate is the basis of nearly all black writing-ink. Ferrous salts give at first only a slight reaction with tannic acid; but the

liquid gradually darkens: characters written with this liquid become quite black in a few hours, and are very permanent.

To an aqueous solution of tannic acid add solution of tartar-emetic; tannate of antimony is precipitated. This reaction and that with gelatine are useful in the quantitative estimation of the amount of tannic acid in various substances.

Tannic acid as it occurs in oak-bark is said to be a glucoside, that is, like several other substances, yields glucose (grape-sugar) when boiled with dilute sulphuric or hydrochloric acid, the other product being gallic acid:—



Catechu (*Catechu pallidum*, B. P.), Gambier, or *Terra Japonica*, an extract of the *Uncaria Gambier*; as well as the true *Catechu*, *Cutch*, or *Terra Japonica*, an extract from the *Acacia Catechu* (*Catechu nigrum*, P. I.) and *A. Suma*; *East-Indian Kino* (*Kino*, B. P.), from the *Pterocarpus marsupium*; also *Bengal or Butea Kino*, from the *Palas* or *Dhak* tree, *Butea frondosa* (*Butea gummi vel Kino Bengalensis*, P. I.); *Elm-bark* (*Ulm Cortex*, B. P.); and some other vegetable products, contain a variety of tannic acid (*mimotannic acid*), which gives a greenish precipitate with neutral solutions of ferric salts. According to Paul and Kingzett it yields, when decomposed, unfermentable sugar, and an acid different to ordinary gallic acid. *Catechu* and Gambier also contain *catechuic acid* or *catechin*, $\text{C}_{13}\text{H}_{12}\text{O}_5$, a body occurring in minute colourless acicular crystals, and, like mimotannic acid, affording a green precipitate with ferric salts.

Bael fruit (*Bela Fructus*, B. P.), from the *Ægle Marmelos*, is said to owe its power as a remedy for dysentery and diarrhoea to a variety of tannic acid; but this is questionable. The rind of the fruit of the *pomegranate* (*Punica granatum*) (*Granati Cortex*, P. I.) contains tannic acid. The astringency of *Pomegranate-root Bark* (*Granati Radicis Cortex*, B. P. and P. I.) is due to a tannic acid (its anthelmintic properties probably to a resinoid matter, or possibly to what Tanret states to be a liquid alkaloid, *pelletierine*, $\text{C}_{16}\text{H}_{30}\text{N}_2\text{O}_2$). A tannic acid also probably gives the astringency to *Logwood* (*Hæmatoxyli Lignum*, B. P.), the colour of which is due to oxidized *hæmatoxylin*. *Rhatany-root bark* (*Krameria Radix*, B. P.)

contains about 20 per cent. of tannic acid, its active astringent principle; rhubarb-root about 9 per cent. *Bear-berry-leaves* (*Uvae Ursi Folia*, B. P.) owe most of their therapeutic power to about 35 per cent. of tannic acid. (The cause of their influence on the kidneys is not yet traced.) They also contain *arbutin*, a crystalline glucoside. *Larch-bark* (*Laricis Cortex*, B. P.), the inner bark of *Pinus larix* or *Larix europæa*, contains, according to Stenhouse, a considerable amount of a tannic acid giving olive-green precipitates with salts of iron, and *larixin* and *larixinic acid* ($C_{10}H_{10}O_6$), a somewhat bitter substance. *Areca nuts* or *Betel nuts* (*Areca*, B. P.), from the *Areca Palm* (*Areca catechu*), contain, according to Flückiger and Hanbury, about 15 per cent. of "tannic matter." The extract of the fruit of *Gab* or *Diospyros embryopteris* (*Diospyri Fructus*, P. I.) is a powerful astringent containing tannic acid. The rhizome of *Geranium maculatum*, *Spotted Orangesbill*, or *Alum-root*, contains both tannic and gallic acids.

Gallic Acid ($H_3C_7H_3O_6$, H_2O) (*Acidum Gallicum*, B. P.) occurs in small quantity in oak-galls and other vegetable substances, but is always prepared from tannic acid. Powdered galls are moistened with water and set aside in a warm place for five or six weeks, or until a little treated with water and filtered yields a solution which is only slightly precipitated by solution of isinglass, occasionally being remoistened; fermentation occurs, and impure gallic acid is formed. The product is treated with about three times its weight of water, boiled to dissolve the gallic acid, filtered, the solution set aside to cool, deposited gallic acid collected, drained, pressed between folds of paper to remove all mother-liquor, and, if necessary, purified by recrystallization from water, or by solution in hot water and treatment with animal charcoal, which absorbs colouring-matter. On filtering and cooling, most of the acid separates in the form of fawn-coloured slender acicular crystals. Gallic acid is soluble in about 100 times its weight of cold or three of boiling water, freely in spirit, sparingly in ether.

The nature of the action by which gallic acid is thus pro-

duced is probably similar to that of the action of dilute acids on tannic acid. During the process oxygen is absorbed and carbonic acid gas evolved, the sugar being thus broken up or perhaps prevented from being formed.

Test.—To an aqueous solution of gallic acid add a neutral solution of a ferric salt; a bluish-black precipitate of gallate of iron falls, similar in appearance to tannate of iron. Ferrous salts also are blackened by gallic acid. To more of the solution add an aqueous solution of gelatine; no precipitate occurs. By the latter test gallic is distinguished from tannic acid.

Pyrogallic acid ($C_6H_6O_3$).—This substance sublimes in light feathery crystals when gallic acid is heated. To an aqueous solution add a neutral solution of a ferric salt; a red colour is produced. To another portion add a ferrous salt; a deep-blue colour results.

Test for the three acids.—To three separate small quantities of milk of lime in test-tubes add, respectively, tannic, gallic, and pyrogallic acids; the first slowly turns brown, the second more rapidly, while the pyrogallic mixture at once assumes a beautiful purplish-red colour changing to brown. These reactions are highly characteristic; they are accompanied by absorption of oxygen from the air.

Use of Pyrogallic Acid in Gas-analysis.—A mixture of pyrogallic acid and solution of potash absorbs oxygen with such rapidity and completeness that a strong solution of each, passed up successively by a pipette into a graduated tube containing air or other gas, form an excellent means of estimating free oxygen. The value of this method may be roughly proved by pouring a small quantity of each solution into a phial, immediately and firmly closing its mouth with a cork, thoroughly shaking the mixture, and then removing the cork under water: the water rushes in and occupies about one-fifth of the previous volume of air, indicating that the atmosphere contains one-fifth of its bulk of oxygen. The small amount of carbonic acid gas present in the air is also absorbed by the alkaline liquid; in delicate experiments this should be removed by the alkali before the addition of pyrogallic acid.

URIC ACID ($H_2C_2H_2N_4O_3$) AND OTHER URATES.—Acidulate a

few ounces of human urine with hydrochloric acid, and set aside for twenty-four hours: a few minute crystals of uric acid will be found adhering to the sides and bottom of the vessel and floating on the surface of the liquid.

Microscopical Test.—Remove some of the floating particles by a slip of glass, and examine by a powerful lens or microscope; the chief portion will be found to be in yellowish semi-transparent crystals, more or less square, two of the sides of which are even and two very jagged; but other forms are common. (See the lithographs in the Section on Urinary Sediments.)

Chemical Test.—Collect more of the deposit, place in a watch-glass or small white evaporating-dish, remove adherent moisture by a piece of blotting- or filter-paper, add a drop or two of strong nitric acid, and evaporate to dryness; the residue will be red. When the dish is cold, add a drop of solution of ammonia; a purplish-crimson colour results. The colour is deepened on the addition of a drop of solution of potash.

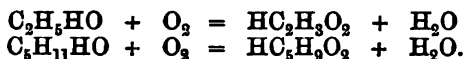
Notes.—Uric acid (or lithic acid) and urates (or lithates) of sodium, potassium, calcium, and ammonium are common constituents of animal excretions. Human urine contains about one part of urate (usually urate of sodium) in 1000. When more than this is present the urate is often deposited as a sediment in the excreted urine, either at once, or after standing a short time. Uric acid or other urate is also occasionally deposited before leaving the bladder, and, slowly accumulating there, forms a common variety of urinary calculus.—Some urates are not definitely crystalline; but when treated with dilute nitric acid or a drop of solution of potash and then a drop or two of acetic acid, jagged microscopic crystals of uric acid are usually formed.—All urates yield the crimson colour when treated as above described.—This colour is due to a definite substance, *murexid* ($C_8H_8N_6O_6$) (from the *murex*, a shell-fish of similar tint); and the test is known as the *murexid test*. The formation of murexid is due to the action of ammonia on *alloxan* ($C_4H_2N_2O_4, 4H_2O$) and other white crystalline products of the oxidation of uric acid

by nitric acid. Murexid is a good dye; it may be prepared from *guano* (the excrement of sea-fowl), which contains a large quantity of urate of ammonium. The excrement of the serpent is almost pure ammonium urate.

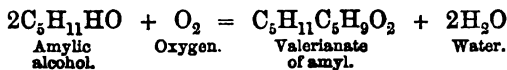
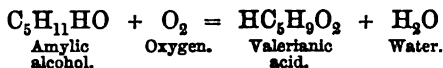
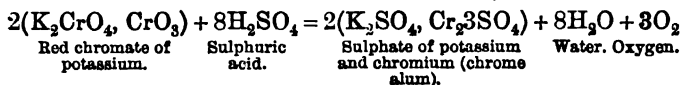
Uric acid and the urates will be again alluded to in connection with the subject of morbid urine.

VALERIANIC ACID or VALERIC ACID ($\text{HC}_5\text{H}_9\text{O}_2$) AND OTHER VALERIANATES.—In a test-tube place a few drops of amylic alcohol (fousel-oil) with a little dilute sulphuric acid and a grain or two of red chromate of potassium, cork the tube, set aside for a few hours, and then heat the mixture; valerianic acid, of characteristic valerian-like odour, is evolved.

Valerianic acid occurs naturally in valerian-root in association with the essential oil from which it is derived (*vide* Index), but is usually prepared artificially, by the foregoing process, from amylic alcohol, to which it bears the same relation as acetic acid does to common alcohol:—

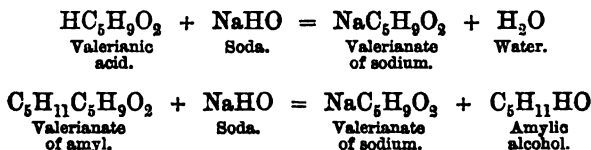


Valerianate of Sodium ($\text{NaC}_5\text{H}_9\text{O}_2$) (*Sodæ Valerianas*, B. P.) is prepared from the valerianic acid and valerianate of amyl obtained on distilling the mixture of amylic alcohol (4 fl. oz.) sulphuric acid ($6\frac{1}{2}$ fl. oz. with 10 of water), and red chromate of potassium (9 oz. in 70 of water). The mixture should stand for several hours before the heat is applied.



The distillate (70 or 80 oz.) is saturated with soda, which not only yields valerianate of sodium with the free valerianic

acid, but decomposes the valerianate of amyl produced at the same time, more valerianate of sodium being formed and some amylic alcohol set free, according to the following equations:—



From the solution of valerianate of sodium (which should be made neutral to test-paper by careful addition of soda solution) the solid white salt is obtained by evaporation to dryness and cautious fusion of the residue. The mass obtained on cooling should be broken up and kept in a well-closed bottle. It is entirely soluble in spirit.

Other Valerianates, as valerianate of zinc (*Zinci Valerianas*, B. P.) and ferric valerianate, may be made by double decomposition of valerianate of sodium with the sulphate or other salt of the metal the valerianate of which is desired, the new valerianate either precipitating or crystallizing out. A hot solution of sulphate of zinc ($5\frac{3}{4}$ parts) and valerianate of sodium (5 parts) in water (40 parts) gives a crop of crystals of valerianate of zinc on cooling.

Tests.—Heated with diluted sulphuric acid, valerianates of the metals give valerianic acid, which has a highly characteristic smell.

The amylic alcohol ($\text{C}_5\text{H}_{11}\text{HO}$) from which valerianates are prepared may contain the next lower homologue, *butylic alcohol* ($\text{C}_4\text{H}_9\text{HO}$). This, during oxidation, will be converted into *butyric acid* ($\text{HC}_4\text{H}_7\text{O}_2$), the next lower homologue of valerianic acid ($\text{HC}_5\text{H}_9\text{O}_2$), and hence the various valerianates be contaminated by some *butyrates*. These are detected by distillation with diluted sulphuric acid and addition of solution of acetate of copper to the distillate, which at once becomes turbid if butyric acid be present. In this reaction valerianic and butyric acids are produced by double decomposition of

the valerianate and butyrate by the sulphuric acid, and distil over on the application of heat. On the addition of acetate of copper ($\text{Cu}_2\text{C}_2\text{H}_3\text{O}_2$) butyrate of copper ($\text{Cu}_2\text{C}_4\text{H}_7\text{O}_2, \text{H}_2\text{O}$) is formed, and, being almost insoluble in water, is at once precipitated, or remains suspended, giving a bluish-white opalescent liquid. Valerianate of copper ($\text{Cu}_2\text{C}_6\text{H}_9\text{O}_2$) is also formed after some time, but is far more soluble than the butyrate, and only slowly collects in the form of greenish oily drops, which gradually pass into greenish-blue hydrous crystalline valerianate of copper (Larocque and Huralt).

VANILLIC ACID ($\text{HC}_8\text{H}_7\text{O}_3$) or VANILLIN ($\text{C}_8\text{H}_8\text{O}_3$) or METHYL-PROTocatechuic ALDEHYD ($\text{C}_7\text{H}_6\text{CH}_3\text{O}_3$); the body to which is due the odour and flavour of vanilla.—The white crystals commonly found on vanilla (the prepared unripe pods of *Vanilla planifolia*), previously termed *vanillin*, were found by Carles to be a weak acid. It occurs in vanilla to the extent of from $1\frac{1}{2}$ to 3 per cent. Vanillin has recently been prepared artificially by Tiemann and Haarman from coniferin, a glucoside existing in the sapwood of pines. The body remaining after the removal of glucose from coniferin, or, indeed, coniferin itself, by action of a mixture of red chromate of potassium and sulphuric acid, yields the vanillin. It also may be obtained by a series of reactions starting from that of carbonic acid on carbolate of potassium; also from the eugenol of oil of cloves. By action of hydrochloric acid, vanillin yields chloride of methyl and protocatechuic aldehyd. Such reactions will be better understood when the pupil has studied succeeding sections on what is commonly termed *Organic Chemistry*. Artificial vanillin is less stable than natural vanillin, perhaps because with the latter is associated a preservative resin.

QUESTIONS AND EXERCISES.

617. What is the constitution of nitrites?
618. Mention a test for nitrites in potable waters.
619. Which nitrite is official?
620. Give the names of some natural and artificial silicates.
621. What is "soluble glass"?
622. Distinguish between silica and silicic acid.
623. How are silicates detected?

426 ANALYTICAL DETECTION OF ACIDULOUS RADICALS.

- 624. What is the quantivalence of silicon?
- 625. Mention the sources, formulæ, and analytical reactions of succinates.
- 626. State the mode of manufacture and tests of sulphocyanates.
- 627. What proportion of tannic acid is contained in galls?
- 628. Describe the official process for the preparation of Tannic Acid.
- 629. Explain the chemistry of "tanning."
- 630. Enumerate the tests for tannic acid.
- 631. What is the assumed constitution of tannic acid?
- 632. Mention official substances other than galls whose astringency is due to tannic acid.
- 633. How is gallic acid prepared?
- 634. By what reaction is gallic distinguished from tannic acid?
- 635. Mention the characteristic properties of pyrogallic acid.
- 636. Explain the murexid test for uric acid.
- 637. Describe the artificial preparation of valerianic acid and other valerianates, giving diagrams or equations.
- 638. What is the formula of valerianic acid?
- 639. How are butyrates detected in presence of valerianates?

DETECTION OF THE ACIDULOUS RADICALS OF SALTS SOLUBLE IN WATER.

Analytical operations may now be resumed, the detection of acidulous radicals being practised for two or three days, and then full analyses made, both for basylous and acidulous radicals. To this end a few compounds of stated metals (potassium, sodium, or ammonium) should be placed in the hands of the practical student for examination according to the following paragraphs and Tables. Mixtures in which both basylous and acidulous radicals may be sought should then be analysed.

In examining salts soluble in water, and concerning which no general information is obtainable, search must first be made for any basylous radicals by the appropriate methods (*vide* pp. 253 or 295-8). Certain metals having been thus detected, a little reflection on the character of their salts will at once indicate what acidulous radicals may be, and what cannot be,

present. Thus, for instance, if the substance under examination is freely soluble in water, and lead is found, only the nitric and acetic radicals need be sought, none other of the lead salts than nitrate or acetate being freely soluble in water. Moreover the salt is more likely to be acetate than nitrate of lead, for two reasons: the former is more soluble than the latter, and is by far the commoner salt of the two. Medical and pharmaceutical students have probably, in dispensing, already learnt much concerning the solubility of salts, and whether a salt is rarely employed or in common use. And although but little dependence can be placed on the chances of a salt being present or absent according to its rarity, still the point may have its proper weight. If, in a mixture of salts, ammonium, potassium, and magnesium have been found associated with the sulphuric, nitric, and hydrochloric radicals, and we are asked how we suppose these bodies may exist in the mixture, it is far more in accordance with common sense to suggest that sal-ammoniac, nitre, and epsom salt were originally mixed together than to suppose any other possible combination. Such appeals to experience regarding the solubility or rarity of salts cannot be made by any one not previously acquainted, or insufficiently acquainted, with the characters of salts; in such cases the relation of a salt to water and acids can be ascertained by referring to the following Table (p. 430) of the solubility or insolubility of about five hundred of the common and rarer salts met with in chemical operations.

The opposite course to the above (namely, to ascertain what acidulous radicals are present in a mixture, and then to appeal to experience to tell what basylous radicals may be and what cannot be present) is impracticable; for acidulous radicals cannot be separated out, one after the other, from one and the same quantity of substance by a similar treatment to that already given for basylous radicals. Indeed such a sifting of acidulous radicals could scarcely be accomplished at all, or only by a vast deal of labour. The basylous radicals must therefore be first detected.

Even when the basylous radicals have been found, the acidulous radicals which may be present must be sought for singly, the only additional aid which can be brought in being the action of sulphuric acid, a barium salt, a calcium salt, nitrate of silver, and ferric chloride on *separate* small portions of the solution under examination, as detailed in the second of the following Tables.

Commence the analysis of an aqueous solution of a salt or salts, the basylous radicals in which are known, by writing out a list of the acidulous radicals which may be, or, if more convenient, of those which cannot be present. To this end consult the following Table (p. 430) of the solubility of salts in water. Look for the name of the metal of the salt in the vertical column; the letters S and I indicate which salts are soluble and which insoluble in water, an asterisk attached to the S meaning that the salt is slightly soluble. The acidulous part of the name is given in the top line of the table. All the names are in alphabetical order, for facility of reference.

Some of the salts marked as insoluble in water are soluble in aqueous solutions of soluble salts, a few forming soluble double salts. To characterize salts as soluble, slightly soluble, or insoluble, only roughly indicates their relation to water: on the one hand, very few salts are absolutely insoluble in water; on the other, there is a limit to the solubility of every salt.

If only one, two, or perhaps three given acidulous radicals can be in the liquid, test directly for it or them according to the reactions given in the previous pages. If several may be present, pour small portions of the solution, rendered neutral if necessary by ammonia, into five test-tubes, and add respectively sulphuric acid, nitrate or chloride of barium, chloride of calcium, nitrate of silver, and ferric chloride; then consult the Table on page 431, in order to correctly interpret the effects these reagents may have produced.

REMARKS ON THE TABLE, page 431.

The first point of value to be noticed in connection with this Table is one of a negative character; namely, if either of the reagents gives no reaction it is self-evident that the salts which it decomposes with production of a precipitate must be absent. Then, again, if the action of one of the reagents indicates the absence of certain acidulous radicals, those radicals cannot be precipitated by the other reagents; thus, if the action of sulphuric acid points to the absence of sulphides, sulphites, carbonates, cyanides, and acetates, these salts may

be struck out of the other lists, and the examination of subsequent precipitates be so far simplified. Or, if the barium precipitate is soluble in hydrochloric acid and the calcium precipitate in acetic acid, neither sulphates nor oxalates can be present. Observing these and other points of difference, which will be seen on careful and thoughtful reflection, and remembering the facts suggested by a knowledge of what basylous radicals are present, one acidulous radical after the other may be struck off as absent or present, leaving only one or two as the objects of special experiment. Among the chief difficulties to be encountered will be the separation from each other of chlorides, bromides, iodides, and cyanides, or of tartrates from citrates, and confirmatory tests of the presence of certain compounds. These may all be surmounted on referring back to the reactions of the various radicals, as described under their hydrogen salts, the acids.

In rendering a solution neutral, for the application of the various group-tests, the employment of any large amount of acid or of alkali must be noted, the presence of actual alkalies (that is *hydrates*) or of acids, respectively, being thereby indicated.

Sulphuric acid, the first group-test, may itself yield, especially when heated with some solid substances, sulphurous acid or hydrosulphuric acid (*see pp. 315 and 317*); hence the production of the latter acids from a *diluted* solution only is evidence of the presence of a sulphide or sulphite.

In the precipitate produced by chloride of barium, the second group-test, the oxalic radical may be specially sought by the test described in the "note" on p. 353.

Chloride of calcium does not precipitate citrates readily or completely in the cold: therefore the mixture should be filtered and the filtrate boiled; calcium citrate then falls. Calcium tartrate is soluble in solution of chloride of ammonium when quite freshly precipitated, but not after it has become crystalline. From their solution in chloride of ammonium, tartrate of calcium is mostly precipitated by ammonia, and citrate on boiling.

The rarer acidulous radicals will very seldom be met with. *Benzoates*, *hippurates* (which give benzoic acid), *hypochlorites*, *hyposulphites*, *nitrites*, and *valerianates* show themselves under the sulphuric treatment. *Ferrocyanides*, *ferridcyanides*, *mecconates*, *succinates*, *sulphocyanates*, *tannates*, and *gallates* appear among the salts whose presence is indicated by ferric chloride;

TABLE OF THE SOLUBILITY OR INSOLUBILITY OF SALTS IN WATER.

	Acetate.	Arseniate.	Arsenite.	Carbonate.	Chloride.	Citrate.	Chromate.	Cyanide.	Hydride.	Iodide.	Nitrate.	Oxalate.	Oxide.	Phosphate.	Sulphate.	Sulphide.	Sulphite.	Tartrate.
Aluminium	.	.	.	.	.	.	.	.	.	.	.	.	.	.	.	.	.	.
Ammonium	S	I	I	I	S	S	S	I	I	S	S	I	I	I	S	I	I	S
Antimony	S	I	I	I	S	S	S	I	I	S	S	I	I	I	S	I	I	S
Barium	S	I	I	I	S	S	S	I	I	S	S	I	I	I	S	I	I	S
Bismuth	S	I	I	I	S	S	S	I	I	S	S	I	I	I	S	I	I	S
Cadmium	S	I	I	I	S	S	S	I	I	S	S	I	I	I	S	I	I	S
Calcium	S	I	I	I	S	S	S	I	I	S	S	I	I	I	S	I	I	S
Chromium	S	I	I	I	S	S	S	I	I	S	S	I	I	I	S	I	I	S
Cobalt	S	I	I	I	S	S	S	I	I	S	S	I	I	I	S	I	I	S
Copper	S	I	I	I	S	S	S	I	I	S	S	I	I	I	S	I	I	S
Ferro	S	I	I	I	S	S	S	I	I	S	S	I	I	I	S	I	I	S
Ferrous	S	I	I	I	S	S	S	I	I	S	S	I	I	I	S	I	I	S
Gold	S	I	I	I	S	S	S	I	I	S	S	I	I	I	S	I	I	S
Lead	S	I	I	I	S	S	S	I	I	S	S	I	I	I	S	I	I	S
Magnesium	S	I	I	I	S	S	S	I	I	S	S	I	I	I	S	I	I	S
Manganese	S	I	I	I	S	S	S	I	I	S	S	I	I	I	S	I	I	S
Mercuric	S	I	I	I	S	S	S	I	I	S	S	I	I	I	S	I	I	S
Mercurous	S	I	I	I	S	S	S	I	I	S	S	I	I	I	S	I	I	S
Nickel	S	I	I	I	S	S	S	I	I	S	S	I	I	I	S	I	I	S
Platinum	S	I	I	I	S	S	S	I	I	S	S	I	I	I	S	I	I	S
Potassium	S	I	I	I	S	S	S	I	I	S	S	I	I	I	S	I	I	S
Silver	S	I	I	I	S	S	S	I	I	S	S	I	I	I	S	I	I	S
Sodium	S	I	I	I	S	S	S	I	I	S	S	I	I	I	S	I	I	S
Stannic	S	I	I	I	S	S	S	I	I	S	S	I	I	I	S	I	I	S
Stannous	S	I	I	I	S	S	S	I	I	S	S	I	I	I	S	I	I	S
Strontium	S	I	I	I	S	S	S	I	I	S	S	I	I	I	S	I	I	S
Zinc	S	I	I	I	S	S	S	I	I	S	S	I	I	I	S	I	I	S

TABLE TO AID IN THE DETECTION OF CHLORIDES, BROMIDES, IODIDES, CYANIDES, NITRATES, CHLORATES, BORATES, ACETATES, SULPHIDES, SULPHITES, SULPHATES, CARBONATES, OXALATES, TARTRATES, PHOSPHATES AND CITRATES, IN A NEUTRAL AQUEOUS SOLUTION.

(For remarks concerning this Table see pages 428, 429 and 432.)

Sulphuria acid decomposes	Chloride of Barium precipitates	Chloride of Calcium precipitates	Nitrate of Silver precipitates	Ferric Chloride precipitates	Not precipitated
Sulphides. Sulphites. Carbonates with effervescence and sulphurous acid gases, known by their smell, and carbonic acid gas, which has no special odour, being evolved. Cyanides, with production of the odour of hydrocyanic acid. Acetates, with production of the odour of acetic acid when the solution is warmed.	Borates. Sulphites. Sulphates. Carbonates. Oxalates. Tartrates. Phosphates. Citrates. Of these white barium precipitates, the sulphate is the only one insoluble in hydrochloric acid; the tartrate and citrate char when heated on platinum foil; the sulphite and carbonate are decomposed with effervescence by acids.	Borates. Oxalates. Sulphites. Tartrates. Sulphates. Phosphates. Carbonates. Citrates. Of these white calcium precipitates, the sulphate only is sol. in much water; the borate, carbonate, and citrate are sol. in solution of chloride of ammonium; all are sol. in acetic acid except oxalate and some tartrate and sulphate; are all sol. in hydrochloric acid, much sulphate excepted; the dry tartrate and citrate char when heated; the sulphite and carbonate effervesce with acids.	Chlorides, white. Bromides, white. Iodides, yellow. Cyanides, white. Borates, white. Sulphides, black. Sulphites, white. Carbonates, white. Oxalates, white. Tartrates, white. Phosphates, yellow. Citrates, white. Of these silver precipitates, the chloride, bromide, iodide, cyanide and sulphide are insoluble in dilute nitric acid; the rest soluble.	Borates, yellowish. Sulphides, black. Carbonates, reddish. Oxalates, yellow. Phosphates, yellow-white. Gives red colour with acetates, if neutral.	Nitrates. Chlorates. Apply special tests.

Note.—The student should precede the examination of aqueous solutions of salts until able to detect acidulous radicals with facility and precision. For this purpose he may finish the analyses of salts or solutions already examined for common and rarer metals, or have aqueous solutions of salts, or the salts themselves, specially prepared for present use, the metals of the salts being stated. He will then be in a position to effectively study the analysis of salts which may or may not be soluble in water, examining them for both basyons and acidulous radicals.

formiates, *hypophosphites*, *malates*, and others by nitrate of silver. *Urates* char when heated, giving an odour resembling that of burnt feathers.

In actual practice the analyst nearly always has some clue to the nature of rarer substances placed in his hands.

If chromium and arsenicum have been detected among the basylous radicals, those elements may be present in the form of *chromates*, *arseniates*, and *arsenites*, yielding with chloride of barium yellow chromate of barium and white arseniate and arsenite of barium, and with nitrate of silver red chromate, brown arseniate, and yellow arsenite of silver.

QUESTIONS AND EXERCISES.

640. In analysing an aqueous solution of salts, for which radicals would you first search, the basylous or the acidulous? and why?

641. In an aqueous solution there have been found magnesium (Mg) and potassium (K), with the sulphuric radical (SO_4), and iodine (I); state the nature of the salts which were originally dissolved in the water, and mention the principles which guide you to the conclusions.

642. Give a sketch of the method by which to analyse a neutral or only faintly acid aqueous liquid for the acidulous radicals of salts. In what stage of the process would the following salts be detected?

- a. Carbonates and Sulphates.
- b. Oxalates.
- c. Tartrates and Nitrates.
- d. Acetates and Sulphites.
- e. Bromides and Cyanides.
- f. Borates.
- g. Iodides and Phosphates.
- h. Chlorates, Oxalates, and Acetates.
- i. Chlorides and Iodides.
- j. Sulphites.
- k. Sulphides, Carbonates, and Nitrates.
- l. Citrates and Sulphates.

643. Nitrate of silver gives no precipitate in an aqueous solution; what acidulous radicals may be present?

644. Chloride of barium gives no precipitate in a neutra solu-

tion, but nitrate of silver a white; what acidulous radicals are indicated?

645. Ferric chloride produces a deep red colour in a solution, chloride of calcium yielding no precipitate; what salts may be present? and how may they be distinguished from each other?

646. Ferric chloride gives a black precipitate in a solution in which sulphuric acid develops no odour; to what is the effect due?

ANALYSIS OF SALTS.

SINGLE OR MIXED, SOLUBLE OR INSOLUBLE.

Thus far all material substances, especially those of pharmaceutical interest, have been regarded as being definite compounds, and as having certain well-defined parts, termed, for convenience, basylous and acidulous respectively: moreover attention has been designedly restricted to those definite compounds which are soluble in water. But there are many substances having no definite or known composition; and of those having definite composition there are many having no definite or ascertained parts. Again, of those having definite composition, and whose constitution admits of the entertainment of theory, there are many insoluble in water.

Chemical substances of whose composition or constitution little or nothing is at present known, are chiefly of animal and vegetable origin, and figure in tables of analyses under the convenient collective title of "extractive matter;" they are not of immediate importance, and may be omitted.

Of substances which are definite in composition, but whose parts or radicals, if they have any, are unknown or imperfectly known, there are only a few (such as the alkaloids, amylaceous and saccharine matters, the glucosides, alcoholic bodies, albumenoid, fatty, resinoid, and colorific substances) which have any considerable amount of medical or pharmaceutical interest; these will be noticed subsequently.

Definite compounds most frequently present themselves; and of these by far the larger proportion (namely, the salts soluble in water) have already been fully studied. There remain, however, many salts which are insoluble in water, but which must be brought into a state of solution before they can be effectively examined from an analytical, pharmaceutical, or a physiological point of view. The next subject

of laboratory work is, therefore, the analysis of substances which may or may not be soluble in water. This will involve no other analytical schemes than those which have been given, will in only one or two cases increase the difficulty of the analysis of a precipitate produced by a group-reagent, but will give roundness, completeness, and a practical bearing to the reader's analytical knowledge. Such a procedure will at the same time bring into notice the methods by which substances insoluble in water are manipulated for pharmaceutical purposes, or made available for use as food by plants, or as food and medicine by man and animals generally.

Preliminary Examination of Solid (chiefly mineral) Salts.

Before attempting to dissolve a salt for analysis, its appearance and other physical properties should be noted, and the influence of heat and strong sulphuric acid be ascertained. If the operator knows how to interpret what is thus observed, and to what extent to place confidence in the observations, he may more certainly obtain a high degree of precision in analysis, and will always gain some valuable negative information. But if he has only slight experience of the appearance and general properties of bodies, or has the habit of turning what should be inferences from tentative processes into foregone conclusions, he should omit the preliminary examination altogether, or only follow it out under the guidance of a judicious tutor; for it is impracticable here to do more than hint at the results which may be obtained by such an examination, or to so adapt descriptions as to prevent a student from allowing unnecessary weight to preconceived ideas.

Whatever be the course pursued, short memoranda describing results should invariably be entered in the note-book.

1. Examine the physical characters of the salt in various ways, but never, or only rarely, by the palate, on account of the danger to be apprehended.

If the salt is white, coloured substances cannot be present; if coloured, the tint may indicate the nature of the substance or of one of its constituents, supposing that the learner is already acquainted with the colours of salts. Closer observation, aided perhaps by a lens, may reveal the occurrence, in a pulverulent mixture, of small crystals or pieces of a single substance; these should be picked out by a needle and ex-

aminated separately. In a powder or roughly divided mixture of substances, the process of sifting (through such sieves as muslin of different degrees of fineness) often mechanically separates substances, and thus greatly facilitates analysis. The body may present an undoubted metallic appearance, in which case only the metals existing under ordinary atmospheric conditions need be sought. Peculiarity in smell reveals the presence of ammonia, hydrocyanic acid, hydrosulphuric acid, etc. Between the fingers a substance is, perhaps, hard, soft, or gritty; consequent inferences follow. Or the matter may be heavy, like the salts of barium or lead; or light, like the carbonates and hydrates of magnesium; or may be one of the pharmaceutically well-known class of "scale" preparations.

2. Place a grain or two of the salt in a small dry test-tube or in a piece of ordinary tubing, closed at one end, and heat it, at first gently, then more strongly, and finally, if necessary, by the blowpipe.

Gases or vapours of characteristic appearance or odour may be evolved; such as iodine, nitrous fumes, sulphurous, hydrocyanic, or ammoniacal gases. Much steam given by a dry substance indicates either hydrates or salts containing water of crystallization. (A small quantity of interstitial moisture often causes heated crystalline substances to *decrepitate*—from *decrepo*, I crackle—that is, break up with slight explosive violence, owing to the expansive force of the steam suddenly generated.) A sublimate may be obtained, due to salts of mercury or arsenicum, to oxalic or benzoic acid, or to sulphur free or as a sulphide—a salt wholly volatile containing such substances only. The compound may blacken, pointing to the presence of organic matter—which, in common definite salts, will probably be in the form of acetates, tartrates, and citrates, or as common salts of the alkaloids morphia, quinia, strychnia, or as starch, sugar, salicin, or in other definite or indefinite forms common in pharmacy, and for which tests will be given in subsequent pages. If no charring occurs, the important fact that no organic matter is present is established. The residue may change colour from presence or development of oxide of zinc, oxide of iron, etc., or melt from the presence of a fusible salt and absence of any large proportion of infusible salt, or be unaltered, showing the absence of any large amount of such substances.

3. Place a grain or two of the salt in a test-tube, add a drop or two of strong sulphuric acid, cautiously smelling any gas that may be evolved; afterwards slowly heat the mixture, noticing the effect, and stopping the experiment when any sulphuric fumes begin to escape.

Iodine, bromine, and nitrous or chlorinoid fumes will reveal themselves by their colour, indicating the presence of iodides, bromides, iodates, bromates, nitrates, and chlorates. The evolution of a colourless gas fuming on coming into contact with air, and having an irritating odour, points to chlorides, fluorides, or nitrates. Gaseous products having a greenish colour and odour of chlorine indicate chlorates, hypochlorites, or chlorides mixed with other substances. Slight sharp explosions betoken chlorates. Evolution of colourless gas may proceed from cyanides, acetates, sulphides, sulphites, carbonates, or oxalates. Charring will be due to citrates, tartrates, or other organic matter. If none of these effects are produced, most of the bodies are absent or only present in minute quantity. The substances apparently unaffected by the treatment are metallic oxides, borates, sulphates, and phosphates.

4. Exposure of the substance to the blowpipe-flame, on platinum wire with or without a bead of borax or of microcosmic salt (phosphate of sodium, ammonium, and hydrogen, NaAmHPO_4)—on platinum foil, in a porcelain crucible, or on a crucible lid, with or without carbonate of sodium—on charcoal, alone or in conjunction with carbonate of sodium, cyanide of potassium, or nitrate of cobalt, will sometimes yield important information, especially to one who has devoted much attention to reactions producible by the blowpipe-flame. The medical or pharmaceutical student, however, will seldom have time to work out this subject to an extent sufficient to make it a trustworthy guide in analysis. (See Plattner and Muspratt "On the Use of the Blowpipe," and a chapter in Galloway's "Manual of Qualitative Analysis.")

Methods of dissolving and analysing single or mixed solid substances.

Having submitted the substance to preliminary examination,

proceed to dissolve and analyse by the following methods. These operations consist in treating a well powdered substance consecutively with cold or hot water, hydrochloric acid, nitric acid, nitrohydrochloric acid, or fusion with alkaline carbonates and solution of the product in water and acid. Resulting liquids are analysed in the manner already described, or by slightly modified processes as detailed in the following paragraphs.

Solution in Water.—Boil about a grain of the salt presented for analysis in about a third of a test-tubeful of water. If it dissolves, prepare a solution of about 20 or 30 grains in half an ounce or more of water, and *proceed with the analysis in the usual way*, testing first for the basylous radical or radicals by the proper group reagents (HCl , H_2S , AmHS , Am_2CO_3 , Am_2HPO_4), pp. 253 or 297, and then for the acidulous radical or radicals, directly or by aid of the prescribed reagents (H_2SO_4 , BaCl_2 , CaCl_2 , AgNO_3 , Fe_2Cl_6), p. 431.

If the salt is not *wholly* dissolved by the water, ascertain whether or not any has entered into solution, by filtering, if necessary, and evaporating a drop or two of the clear liquid to dryness on platinum foil; the presence or absence of a residue gives the information sought. If anything is dissolved prepare a sufficient quantity of solution for analysis and proceed as usual, reserving the insoluble portion of the mixture, after thoroughly exhausting with water, for subsequent treatment by acids.

Solution in Hydrochloric Acid.—If the salt is insoluble in water, digest about a grain of it (or of the insoluble portion of a mixed salt) in a few drops of hydrochloric acid, adding water and boiling if necessary. If the salt wholly dissolves, prepare a sufficient quantity of the liquid, noticing whether or not any effervescence (due to the presence of sulphides, sulphites, carbonates or cyanides) occurs, and proceed with the analysis as before, except that the first step, the addition of hydrochloric acid, may be omitted.

The analysis of this solution will in most respects be simpler than that of an aqueous solution, inasmuch as the majority of salts (all those soluble in water) will be absent. This acid solution will, in short, only contain :—chlorides produced by

the action of the hydrochloric acid on sulphides, sulphites, carbonates, cyanides, oxides, and hydrates; and certain borates, oxalates, phosphates, tartrates, and citrates (possibly silicates and fluorides), which are insoluble in water, but soluble in acids without apparent decomposition. The first four—sulphides, sulphites, carbonates, and cyanides—will have revealed themselves by the occurrence of effervescence during solution; and the presence of oxides and hydrates may often be inferred by the absence of compatible acidulous radicals. The borates, oxalates, phosphates, tartrates, and citrates alluded to will be reprecipitated in the general analysis as soon as the acid of the solution is neutralized; that is, will come down in their original state when ammonia and sulphhydrate of ammonium are added in the usual course. Of these precipitates, only the oxalate of calcium and the phosphates of calcium and magnesium need occupy attention now; for oxalate and phosphate of barium seldom or never occur, and the borates, tartrates, and citrates met with in medicine or in general analysis are all soluble in water. These phosphates and oxalates, then, will be precipitated in the course of analysis along with iron, their presence not interfering with the detection of any other metal. If, from the unusually light colour of the ferric precipitate, phosphates and oxalates are suspected, it is treated according to the following Table (reference to which should be inserted in the Table for metals, under Fe, p. 253, and in the long Table, p. 296).

PRECIPITATE OF PHOSPHATES, OXALATES, AND FERRIC HYDRATE.

Dissolve in HCl, add citric acid, then NH_4HO , and filter.

Filtrate Fe. Add HCl and $\text{K}_4\text{Fcy.}$ Blue ppt.	Precipitate Ca_32PO_4 , CaC_2O_4 , Mg_32PO_4 . Boil in acetic acid and filter.	
	Insoluble CaC_2O_4 . White.	Filtrate Ca_32PO_4 , Mg_32PO_4 . Add $\text{Am}_3\text{C}_2\text{O}_4$, stir, filter.
		Precipitate white, indicating Ca_32PO_4 .
		Filtrate, add AmHO . White ppt. MgNH_4PO_4 .

In analysing phosphates and oxalates advantage is also frequently taken of the facts that the phosphoric radical is wholly removed from solution of phosphates in acid by the addition of an alkaline acetate, ferric chloride, and subsequent ebullition, as described under "Phosphoric Acid" (p. 387), and that dry oxalates are converted into carbonates by heat, as mentioned under "Oxalic Acid" (p. 370). (See also p. 387, 4th Anal. Reac.)

Certain arseniates and arsenites, insoluble in water but soluble in hydrochloric acid, may accompany the above phosphates and oxalates if from any cause hydrosulphuric acid gas has not been previously passed through the solution, or passed for an insufficient length of time.

If the substance insoluble in water does not *wholly* dissolve in hydrochloric acid, ascertain if any has entered into solution, by filtering, if necessary, and evaporating a drop of the clear liquid to dryness on platinum foil; the presence or absence of a residue gives the information sought. If anything is dissolved, prepare a sufficient quantity of solution for analysis and proceed as usual, reserving the insoluble portion of the mixture, after thoroughly exhausting with hydrochloric acid and well washing with water, for the following treatment by nitric acid.

Solution in Nitric Acid.—If the salt is insoluble in water and hydrochloric acid, boil it, (or that part of it which is insoluble in those menstrua) in a few drops of nitric acid. If it wholly dissolves, remove excess of acid by evaporation, dilute with water, and proceed with the analysis.

This nitric solution can contain only a few substances; for nearly all salts soluble in nitric acid are also soluble in hydrochloric acid, and therefore will have been removed previously. Some of the metals, however (Ag, Cu, Hg, Pb, Bi), as well as amalgams and alloys, unaffected or scarcely affected by hydrochloric acid, are readily attacked and dissolved by nitric acid. Many of the sulphides, also, insoluble in hydrochloric acid, are dissolved by nitric acid, usually with separation of sulphur. Calomel is converted, by long boiling with nitric acid, into mercuric chloride and nitrate. The nitrates here produced are soluble in water.

This nitric solution, as well as the hydrochloric and aqueous solutions, should be examined separately. Apparently, time would be saved by mixing the three solutions together and making one analysis. But the object of the analyst is to separate every radical from every other; and when this has been partially accomplished by solvents, it would be unwise to again mix and separate a second time. Moreover solvents often do what the chemical reagents cannot—namely separate *salts* from each other. This is important, inasmuch as the end to be attained in analysis is not only an enumeration of the radicals present, but a statement of the actual condition in which they are present; the analyst must, if possible, state of what salts a given mixture was originally formed—how the basylous and acidulous radicals were originally distributed. In attempting this, much must be left to theoretical considerations; but a process by which the salts themselves are separated is of trustworthy practical assistance; hence the chief advantage of analysing separately the solutions resulting from the action of water and acids on a solid substance.

Solution in Nitro-Hydrochloric Acid.—If the salt or any part of a mixture of salts is insoluble in water, hydrochloric acid, and nitric acid, digest it in nitro-hydrochloric acid, warming, or even boiling gently, if necessary; evaporate to remove excess of acid, dilute, and proceed as before.

Sulphide of mercury and substances only slowly attacked by hydrochloric or nitric acid, as, for example, calomel and ignited ferric oxide, are sufficiently altered by the free chlorine of aqua regia to become soluble.

Analysis of Insoluble substances.

If the substance is insoluble in water and acids, it is one or more of the following substances:—Sand and certain silicates, such as pipeclay and other clays; fluor spar; cryolite (3NaF , AlF_3); sulphates of barium, strontium, and possibly calcium; tinastone; antimonious oxide; glass; felspar (double silicate of aluminium and other metals); chloride, bromide or iodide of silver; sulphate of lead. It may also be or contain carbon or carbonaceous matter, in which case it is black and combustible, burning entirely or partially away when heated in the air—or

be or contain sulphur, in which case sulphurous gas is evolved, detected by its odour, when the substance is heated in air. A drop of solution of sulphhydrate of ammonium added to a little of the powder, will at once indicate the presence or absence of salts of such metals as lead and silver. For the other substances proceed according to the following (Bloxam's) method :—

Four or five grains of the dry substance are intimately mixed with twice the quantity of dried carbonate of sodium, and this mixture well rubbed in a mortar with five times its weight of *deflagrating flux* (1 of finely powdered charcoal to 6 of nitre). The resulting powder is placed in a thin porcelain dish, or crucible, or clean iron tray, and a lighted match applied to the centre of the heap. Deflagration ensues, and decomposition of the various substances occurs, the acidulous radicals going to the alkali metals to form salts soluble in water, the basylous radicals being simultaneously converted into carbonates or oxides. The mass is boiled in water for a few minutes, the mixture filtered, and the residue well washed. The filtrate may then be examined for acidulous radicals and aluminium, and the residue dissolved in dilute hydrochloric acid and analysed by the ordinary method.

The only substance which resists this treatment is chrome iron-ore.

To detect alkali in felspar, glass, or cryolite, Bloxam recommends deflagration of the powdered mineral with one part of sulphur and six of nitrate of barium. The mass is boiled in water, the mixture filtered, hydrate and carbonate of ammonium added to remove barium, the mixture again filtered, and the filtrate evaporated and examined for alkalis by the usual process.

Hydrates and Oxides.

If no acidulous radical can be detected in a substance under analytical examination, or if the amount found is obviously insufficient to saturate the quantity of basylous radical present, the occurrence of oxides or hydrates, or both, may be suspected. Confirmation of their presence will be found in the general rather than in any special behaviour of the substances. Some

hydrates yield water when heated—in a dry test-tube held nearly horizontally in a flame, so that moisture may condense on the cool part of the tube. Some oxides yield oxygen—detected by heating in a test-tube, and inserting the incandescent end of a strip of wood. Soluble hydrates cause abundant evolution of ammonia gas when heated with solution of chloride of ammonium. Soluble hydrates also give characteristic precipitates with the various metallic solutions. Hydrates and oxides insoluble in water not only neutralize much nitric acid or acetic acid, but are thereby converted into salts soluble in water. Most oxides and hydrates have a characteristic appearance. In short, some one or more properties of an oxide or a hydrate will generally betray its presence to the student who not only has knowledge respecting chemical substances, but has cultivated the faculties of observation and perception.

QUESTIONS AND EXERCISES.

647. Describe the preliminary treatment to which a salt may be subjected prior to systematic analysis.

648. Mention substances which might be recognised by smell.

649. Which classes of salts are heavy, and which light?

650. Name some bodies detectable by their colour.

651. What inference may be drawn from the appearance of steam when dry substances are heated?

652. Why do certain crystals decrepitate?

653. If a powder sublimes on being heated, to what classes of compounds may it belong?

654. When heat causes charring, what conclusion is drawn?

655. No change occurring by heat, which substances cannot be present?

656. Give examples of salts which are identified by their reaction with strong sulphuric acid, and by their comportment in the blowpipe-flame, with or without borax or microcosmic salt.

657. What are the solvents usually employed in endeavouring to obtain a substance in a state of solution? and what is the order of their application?

658. Name a few salts which may be present in an aqueous solution.

659. Mention some common compounds insoluble in water, but soluble in hydrochloric acid.

660. What substances are only attacked by nitric acid or nitrohydrochloric acid?

661. At what stage of analysis do arsenites and arseniates show themselves?

662. Sketch out a method for the complete analysis of a liquid suspected to be an aqueous solution of neutral salts.

663. How can earthy phosphates and oxalates with ferric oxide be separated from each other?

664. How would you proceed to analyse an alloy?

665. By what process may substances insoluble in water or acids be analysed?

666. How would you qualitatively analyse glass?

RECAPITULATORY AND OTHER NOTES ON THE CONSTITUTION OF THE DEFINITE CHEMICAL COMPOUNDS COMMONLY TERMED SALTS.

The molecules of a salt contain *radicals*, which may be either elementary or compound (pp. 31 and 65).

Each radical has a definite exchangeable value (p. 132).

The definite exchangeable values of radicals differ in different series of radicals (pp. 133 and 134).

In one and the same molecule of a salt, two or more different atoms of the same element may possess the two distinct functions of being (*a*) a single definite distinct radical, and (*b*) one member of a group of atoms which together form a single definite distinct radical (p. 303).

The relation to each other, either of the elementary or the compound radicals in organic substances or salts, is apparently far more complex than the relation to each other of the elementary or compound radicals in inorganic or mineral salts (pp. 445 to 476).

The properties of salts are regarded as depending on (*a*) the nature, (*b*) the number (p. 480), and (*c*) the position (p. 481), in relation to each other, of the elementary and compound radicals in a molecule.

Dumas, afterwards Laurent, and then Gerhardt, attempted the classification of salts under such *types* as the following:—



The hydrogen type.



The water type.



The ammonia type.

Other chemists have extended the number of such types of salts. Further, by writing the typical formulæ in the above and other manners, a mode of indicating the facts assumed to be dependent on the position of the atoms in a molecule has been sought to be obtained. Finally, the natural development of this train of thought and of practice has produced the *graphic formulæ* of Kékulé, Frankland, and others.

Caution.—The conjectural or theoretic character of our ideas respecting masses of matter being formed of molecules, and molecules of atoms, and that molecules contain radicals consisting of one or more atoms, must never be lost sight of, highly valuable and practically useful though the hypotheses be (pp. 30, 48, 303, 305, 331, 348).

Berthollet's Laws.—"When we cause two salts to react by means of a solvent, if, in the course of double decomposition, a new salt can be produced less soluble than those which we have mixed, this salt will be produced." "When we apply dry heat to two salts, if, by double decomposition, a new salt can be produced more volatile than the salts previously mixed, this salt will be produced."

Malaguti's Law.—When solutions of two different salts are mixed and metathesis occurs, and four salts result, the proportions of the salts to each other are dependent on the strength or intensity of force with which the respective basylous and acidulous radicals are united.

The state of equilibrium just mentioned may be permanent or temporary. The latter condition obtains when one of the salts which may possibly be produced is insoluble, for as soon as precipitation occurs the equilibrium is upset, and is re-established only to be upset again, and so on until from the four salts there result one in solution and one out of solution. This would seem to be the way in which the laws termed Berthollet's work.

THE CHEMISTRY OF CERTAIN SUBSTANCES OF VEGETABLE AND ANIMAL ORIGIN.

Except alcohol and a few acids, the compounds which have hitherto engaged notice have been of mineral origin. But the two other kingdoms of nature, the animal and vegetable, furnish a large number of definite substances. These, indeed, when discovered, were producible only by organized living structures, and were hence termed *organic* compounds.*

A few of these compounds, of common occurrence in pharmacy and possessing prominent characteristics, may now occupy attention; reactions of the alkaloids and some other principles may be performed, and the methods of examining morbid urine be experimentally studied. There will then remain to be studied certain galenical, as distinguished from chemical substances, solid and liquid, which can only be fairly regarded from a pharmacist's rather than a chemist's point of view, and a still larger number, doubtless, not yet brought within the grasp of chemist or pharmacist, and of which, therefore, we must at present be content to remain in ignorance. An opportunity, however, will be afforded of noticing the effect of such organic matter as a vomit or the contents of a stomach, in masking or preventing the reactions by which mineral and vegetable poisons are detected.

ALKALOIDS.

Constitution of Alkaloids or Organic Bases.

Natural alkaloids.—The alkaloids, or alkali-like bodies (εἶδος, *eidos*, likeness), have many analogies with ammonia. Their constitution is not yet known; but they are probably derivatives of a single molecule of ammonia (NH_3), or of double, triple, or quadruple molecules (N_2H_6 , N_3H_9 , N_4H_{12}).

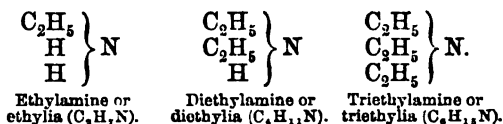
Artificial Alkaloids.—Numerous artificial alkaloids or organic bases, unquestionably having the constitution just mentioned,

* *Organic*, from ὄργανον, *organon*, an organ. A large number of organic compounds can now be obtained artificially—without the aid of a living organism; hence the distinction formerly drawn between organic and inorganic compounds, organic and inorganic chemistry, is fast breaking down.

have already been formed. These are sometimes termed *amines*, and are primary, secondary, and tertiary according as one, two, or three atoms of hydrogen in ammonia have been displaced by radicals, as seen in the following general formulæ (R = any univalent radical. *Vide* Index, "Alcohol radicals") :—



or in the following examples :—



The three classes have also been termed amidogen-, imidogen- and nitrile-bases.

Mode of formation of Artificial Alkaloids.—A few illustrations will suffice. Just as the addition of iodide of hydrogen (HI) to ammonia (that is, the common, tri-hydrogen ammonia, NH_3) gives iodide of common ammonium (NHHHHI or NH_4I), so the addition of iodide of ethyl ($\text{C}_2\text{H}_5\text{I}$ or EtI) (see page 514) to ammonia (NH_3) gives the iodide of ethyl-ammonium (NHHHETI , or NH_3EtI , or $\text{NH}_3\text{C}_2\text{H}_5\text{I}$). A fixed alkali turns out common ammonia (NHHH) from the iodide (or any other salt) of common ammonium; it turns out ethyl-ammonia (NHHET) from the iodide (or any other salt) of ethyl-ammonium. Ethyl-ammonia (or *ethylia*, or ethylamine), NHHET , with iodide of ethyl, EtI , gives iodide of diethyl-ammonium [NHHETEtI , or $\text{NH}_2\text{Et}_2\text{I}$, or $\text{NH}_2(\text{C}_2\text{H}_5)_2\text{I}$]. From the latter, potash turns out diethyl-ammonia (NHEt_2); diethyl-ammonia (*diethylia* or diethylamine) with iodide of ethyl gives iodide of triethyl-ammonium (NHEt_3I). The latter with alkali gives triethyl-ammonia or *triethylia* or triethylamine (NEt_3), and this with iodide of ethyl gives iodide of tetrethyl-ammonium, NEt_4I .

What has just been stated respecting iodide of ethyl is true of other salts of ethyl; and what is true of salts of ethyl is true of salts of an immense number of other radicals—univalent, bivalent, etc.; so that a vast number of artificial alkaloids and their salts can be produced. The reactions are

not always so sharp as those just given. Mixtures of primary, secondary, and tertiary compounds rather than either alone often result in an experiment: but the reactions are typically true.

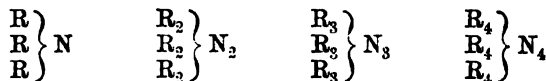
Some of these artificial alkaloids not only resemble natural alkaloids but are strong caustic liquids, like solution of ammonia.

Then, the displacing radical in an artificial alkaloid or its salt may not be of one kind, as indicated in the preceding paragraphs, but of different kinds; and while the radical displacing one atom of hydrogen is keeping its place, any of the many known radicals may occupy the position of one or all of the other atoms of hydrogen. Thus, for example, we have methyl-ethyl-amylamine ($C_8H_{19}N$, or $NCH_3C_2H_5C_5H_{11}$, or $NMeEtAy$), a colourless, oily body, of agreeable aromatic odour. The empirical formulæ of morphia, quinia, etc., may some day be similarly resolvable into rational formulæ. Their artificial production will then quickly follow.

Methylamine (CH_3HHN) and *trimethylamine* ($(CH_3)_3N$), are artificial alkaloids which have been found by Schmidt, in *Mercurialis annua* and *M. perennis*, and previously by Reichardt, who termed it *mercurialine*. Trimethylamine is also produced in large quantities in the dry distillation of the evaporated residue of the spent wash produced in beet root spirit distilleries.

Propylamine or *tritylia* (C_3H_7HHN) is a volatile oil, one product of the destructive distillation of bones and other animal matters.

The organic bases derived from one molecule of ammonia are termed monamines; from two molecules, diamines; from three, triamines; and from four tetramines:—



In these amines, any bivalent, trivalent, or quadrivalent radical may occupy the place of two, three, or four univalent radicals.

Evidence of Constitution of the Natural Alkaloids—Attempts to form artificially the important natural organic bases have hitherto failed; but the primary, secondary, or tertiary character of some of them has been indicated by the intro-

duction or elimination of methyl, ethyl, and other radicals for hydrogen.

Notes on Nomenclature of Natural Alkaloids.—The first syllables of the names of the natural alkaloids recall the name of the plant whence they were obtained, or some characteristic property. It is to be regretted that the last syllable is not either *ine* or *ia*, instead of sometimes one and sometimes the other: general usage seems to be in favour of the latter, a plan that distinguishes the alkaloids from some other substances the names of which end in *ine*, as chlorine, bromine, iodine, fluorine, glycerine, gelatine, etc. The names of the salts of the alkaloids are given on the assumption that the acid unites with the alkaloid without decomposition. Thus hydrochlorate of morphia is regarded as morphia with hydrochloric acid, just as we might assume sal-ammoniac to be ammonia (NH_3) with hydrochloric acid (HCl), and name it hydrochlorate of ammonia (NH_3HCl) instead of chloride of ammonium (NH_4Cl). All acids, even sulphydric, unite with alkaloids and form salts having similar names.

Antidotes.—In cases of poisoning by alkaloids, emetics and the stomach-pump must be relied on rather than chemical agents. But astringent liquids may be administered, for tannic acid precipitates many of the alkaloids from their aqueous solution, absorption of the poison being thus possibly retarded.

MORPHIA, OR MORPHINE.

Formula $\text{C}_{17}\text{H}_{19}\text{NO}_3$, H_2O . Molecular weight 303.

Occurrence.—Morphia occurs in opium (the inspissated juice of the fruit, *Papaveris capsulæ*, of the White Poppy, *Papaver somniferum*) as meconate of morphia [$(\text{C}_{17}\text{H}_{19}\text{NO}_3)_2$, $\text{C}_7\text{H}_4\text{O}_7$, $5\text{H}_2\text{O}$; Dott]. The dried poppy-capsule of pharmacy contains opium principles, but varying much in nature and proportion. Ordinary Asia-Minor opium (Turkey, Smyrna, or Constantinople opium) contains, when dried, from 10 to 15 per cent. of morphia.

Process for Hydrochlorate.—The hydrochlorate, $\text{C}_{17}\text{H}_{19}\text{NO}_3$, HCl , $3\text{H}_2\text{O}$ (*Morphiæ Hydrochloras*, B. P.), occurs in slender white acicular crystals; it is prepared by simply decomposing an aqueous infusion of opium with chloride of calcium, meconate of calcium and hydrochlorate of morphia being produced. (If the infusion, which is always acid, be first nearly neutral-

ized by the cautious addition of small quantities of a very dilute solution of ammonia, the chloride of calcium then at once causes a precipitate of meconate of calcium, which can be filtered off, leaving a coloured solution of hydrochlorate of morphia. On the large scale—*vide* B. P.—the details are somewhat different.) The salt is partially purified by crystallization from the evaporated liquid, then by treatment of the solution of the impure hydrochlorate by animal charcoal, and lastly by precipitation of the morphia from the still coloured liquid by ammonia and resolution of the morphia in hot dilute hydrochloric acid; hydrochlorate of morphia separates out on cooling.

Hydrochlorate of morphia deposited from a hot solution in about twenty times its weight of alcohol is anhydrous.

Morphia may also, of course, be prepared by the methods given for its quantitative separation from opium (*see* Index).

Process for Acetate.—Acetate of morphia ($C_{17}H_{19}NO_3C_2H_4O_2$) (*Morphiæ Acetas*, B. P.) is a white pulverulent salt prepared by dissolving morphia in acetic acid, the morphia being prepared from a solution of the hydrochlorate by precipitation with ammonia. Eight parts of hydrochlorate yield about seven of acetate. One grain of acetate, so made, in twelve minims of water forms the *Injectio Morphiæ Hypodermica*, B. P.

Both the hydrochlorate and acetate of morphia are soluble in water, but the solution is not stable unless acidulated and containing alcohol; hence the official solutions, 4 grains in one ounce (*Liquor Morphiæ Hydrochloratis*, B. P., and *Liquor Morphiæ Acetatis*, B. P.), consist of three parts water and one part rectified spirit, a few minims per ounce of hydrochloric or acetic acid being added. Even solid acetate of morphia is unstable, slowly dissociating into acetic acid and morphia, hence the acid odour of acetate of morphia. The other preparations official in the British Pharmacopœia are *Suppositoria Morphiæ*, *Trochisci Morphiæ*, and *Trochisci Morphæ et Ipecacuanhæ*.

When a simple but strong solution of acetate of morphia in water is required (for hypodermic injection), one part of the salt, if recently made, may be dissolved in six parts of water (1 grain in 11 grains, B. P.). If the acetate of morphia is old, it either will not dissolve, or, if dissolved by aid of warmth and a little acetic acid, will soon be re-deposited. The aqueous solution, however well made, cannot be kept in its normal condition for any great length of time.

Other alkaloids exist in opium. In the above process a considerable quantity of an alkaloid of very weak basic properties, *narcotine* ($C_{22}H_{23}NO_7$) (*Narcotina*, P. I.), remains in the exhausted opium, and may be extracted by digesting in acetic acid, filtering, and precipitating by ammonia. It crystallizes in brilliant needles from alcohol or ether. By oxidation it yields *cotarnine* and an acid termed *opianic*. *Codeia* ($C_{18}H_{21}NO_3$, H_2O) is soluble in the slight excess of ammonia employed in precipitating the morphia. From the mother-liquors there have also been obtained *thebaia* ($C_{19}H_{21}NO_3$), *papaverine* ($C_{21}H_{21}NO_4$), *opianine* ($C_{21}H_{21}NO_7$?), *narceia* ($C_{23}H_{29}NO_9$), *cryptopia* ($C_{31}H_{23}NO_5$), *meconine* ($C_{10}H_{10}O_4$), *meconiosine* ($C_8H_{10}O_2$), *laudanine* ($C_{20}H_{25}NO_4$), *codamine* ($C_{30}H_{25}NO_4$), *gnoscopine* ($C_{34}H_{36}N_2O_{11}$), *pseudomorphia* ($C_{17}H_{19}NO_4$), *protopine* ($C_{20}H_{19}NO_5$), *laudanosine* ($C_{21}H_{27}NO_4$), *hydrocotarnine* ($C_{12}H_{15}NO_3$), *rhœadine* ($C_{30}H_{21}NO_6$), *meconidine* ($C_{31}H_{23}NO_4$), *lanthopine* ($C_{23}H_{25}NO_4$).

A little acetic acid also exists in all opium (D. Brown).

ANALYTICAL REACTIONS.

First Analytical Reaction.—To a minute fragment of a salt of morphia add one drop of water, and warm the mixture until the salt dissolves, then stir the liquid with a glass rod moistened by a strong *neutral* solution of perchloride of iron; a dirty blue colour is produced. This effect is not observed in dilute solutions.

Second Analytical Reaction.—To a drop or two of a strong solution of a morphia salt in a test-tube add a minute fragment of iodic acid (HIO_3 , p. 344); iodine is set free. Into the upper part of the tube insert a glass rod covered with mucilage of starch, and warm the solution; dark-blue starch iodide is produced. If the mixture of morphia and iodic acid be shaken up with chloroform or bisulphide of carbon a violet solution is obtained.

This reaction is only confirmatory of others, as albuminous matters also reduce iodic acid.

Third Analytical Reaction.—To a few drops of an aqueous infusion of opium add a drop of *neutral* solution of perchlorid

of iron; a red solution of meconate of iron is produced. Add solution of corrosive sublimate; the colour is not destroyed. (The red sulphocyanide is decolourized.) Boil; the colour is not discharged. (The red acetate is decomposed by ebullition.)

In cases of poisoning by a preparation of opium, this test is almost as conclusive as a direct reaction of morphia (the poison itself), meconic acid being obtainable from opium only.

Other Reactions.—Add carbonate of sodium to a solution of a salt of morphia; a white precipitate of morphia falls, slowly and of a crystalline character if the solution is dilute. Collect this precipitate and moisten it with neutral solution of perchloride of iron; the bluish tint above referred to is produced.—Add an alkali to a solution of hydrochlorate or acetate of the alkaloid; morphia is precipitated, soluble in excess of the fixed alkali, far less readily so in ammonia.—Moisten a particle of a morphia salt with nitric acid; an orange-red coloration is produced.—To morphia add strong sulphuric acid, mix, and strew nitrate of bismuth on the fluid; the mixture turns dark brown or black.—Heat morphia on platinum foil; it burns entirely away.

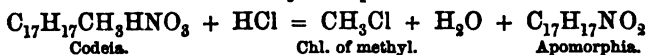
APOMORPHIA ($C_{17}H_{17}NO_2$).

Apomorphia (*ἀπό*, *apo*, from, and *morphia*) is an alkaloid obtained from morphia by Matthiessen and Wright. It possesses remarkable physiological effects; one-tenth of a grain (in aqueous solution) injected under the skin, or a quarter of a grain taken into the stomach, is said to produce vomiting in from four to ten minutes.

Process.—Hydrochlorate of morphia is hermetically sealed in a thick tube with considerable excess of hydrochloric acid, and heated to nearly 300° F. for two or three hours. The product is purified by diluting the contents of the tube with water, precipitating with bicarbonate of sodium, and treating the precipitate with ether or chloroform.—On shaking up the ethereal or chloroform solution with a very small quantity of strong hydrochloric acid, the sides of the vessel become covered with crystals of the hydrochlorate of the new base. These may be drained from the mother-liquor, washed with a

little cold water, in which the salt is sparingly soluble, recrystallized from hot water, and dried on bibulous paper or over sulphuric acid. The formula ($C_{17}H_{17}NO_3, HCl$) indicates that the new alkaloid is derived from morphia by abstraction of the elements of water.

Codeia also, according to the same chemists, yields apomorphia by similar treatment, a reaction that would seem to indicate that codeia is methyl-morphia:—

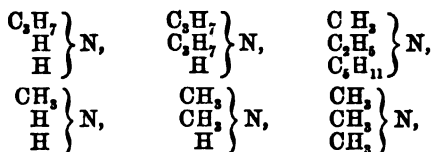


Dr. C. R. A. Wright has recently obtained several new derivatives of codeia.

QUESTIONS AND EXERCISES.

667. Write some general formulæ of artificial alkaloids.

668. Name the substances represented by the following formulæ:—



669. What is the assumed constitution of the salts of the alkaloids?

670. Describe the treatment in cases of poisoning by alkaloids.

671. Give the official process for the preparation of Hydrochlorate of Morphia. In what form does morphia occur in opium?

672. How is Acetate of Morphia prepared?

673. What plan is adopted for preventing the decomposition of the official solutions of morphia?

674. Mention the analytical reactions of morphia.

675. In addition to the reactions of morphia, what test may be employed in searching for opium in a liquid or semifluid material?

676. How is apomorphia prepared? and what are its properties?

677. Describe the relation of morphia to codeia.

QUINIA, OR QUININE.

Formula $C_{20}H_{24}N_2O_2, 3H_2O$. Molecular weight 378.

Source.—Quinia and other similar alkaloids exist in cinchona-bark as kinates. In the yellow bark (*Cinchona Flavæ Cortex*, B. P., from *Cinchona Calisaya*) chiefly quinia is present; in the pale (Loxa or Crown) bark (*Cinchona Pallidæ Cortex*, B. P., chiefly from *C. officinalis*) other alkaloids are more frequently found; while in the red bark (*Cinchona Rubræ Cortex*, B. P., from *C. succirubra*) the alkaloids occur in irregular proportions.

Extraction of the Mixed Alkaloids.—Take 750 grains of finely powdered bark. Make it into a paste with milk of lime (slaked lime about 400 grains and water about 4 ounces). Dry the mixture over a water-bath. Powder the residue and place the whole in a cylindrical percolator. Pour in $3\frac{1}{2}$ fluid ounces of chloroform. When, after standing, packing is complete, allow percolation to commence and to proceed slowly. After a time pour $3\frac{1}{2}$ fluid ounces more chloroform into the percolator. When percolation has ceased transfer the percolate to a retort, and add nearly half an ounce of water and enough dilute sulphuric acid to make the mixture acid to test paper. Recover the chloroform by distilling from a water-bath, and allow the residue to cool. Filter. To the filtrate, which contains the alkaloids as acid sulphates, add ammonia in slight excess. Collect the precipitated alkaloids on a filter, wash, and dry in the air or over a dish of sulphuric acid covered by a bell-glass. For the separation of alkaloids, see Index, "De Vrij's process"; an operation which should not be attempted at this stage of study.

Process for Sulphate.—Sulphate of quinia (*Quinæ Sulphas*, B. P.) is prepared by treating the yellow bark with dilute hydrochloric acid, precipitating the resulting solution of hydrochlorate of quinia by soda, and redissolving the precipitated quinia in the proper proportion of hot dilute sulphuric acid. This, the common commercial sulphate, crystallizes out on cooling in silky acicular crystals, one molecule containing two atoms of quinia ($2C_{20}H_{24}N_2O_2$), one of sulphuric acid (H_2SO_4), and eight of water of crystallization ($8H_2O$).

The details of the official process are as follows :—

Take of

Yellow Cinchona-bark, in coarse powder	1 pound,
Hydrochloric Acid	3 fluid ounces,
Distilled Water	a sufficiency,
Solution of Soda	4 pints (or less),
Diluted Sulphuric Acid	a sufficiency.

Dilute the hydrochloric acid with ten pints of the water. Place the cinchona-bark in a porcelain basin, and add to it as much of the diluted hydrochloric acid as will render it thoroughly moist. After maceration, with occasional stirring, for twenty-four hours, place the bark in a displacement-apparatus, and percolate with the diluted hydrochloric acid until the solution which drops through is nearly destitute of bitter taste. Into this liquid (solution of hydrochlorate of quinia) pour some solution of soda, agitate well, let the precipitate (quinia) completely subside, decant the supernatant fluid, collect the precipitate on a filter, and wash it with cold distilled water until the washings cease to have colour. Transfer the precipitate to a porcelain dish containing a pint of distilled water, and, applying to this the heat of a water-bath, gradually add diluted sulphuric acid until very near the whole of the precipitate has been dissolved and a neutral liquid has been obtained. (Or add about *half* the precipitated quinia to some water in an evaporating-basin, warm the mixture and pour in diluted sulphuric acid until the precipitate has dissolved and the liquid is neutral or only faintly acid, then add the other half, stir well, and again heat the liquid.) Filter the solution (sulphate of quinia) while hot through paper, wash the filter with boiling distilled water, concentrate till a film forms on the surface of the solution, and set it aside to crystallize. The crystals should be dried on filtering-paper without the application of heat.

Sulphate of quinia, the *common* or so-called *disulphate*— $(C_{20}H_{34}N_2O_2)_2, H_2SO_4, 8H_2O$,—is only slightly soluble in

water; on the addition of dilute sulphuric acid the so-called *neutral sulphate*, or *soluble sulphate* ($C_{30}H_{24}N_2O_9, H_2SO_4, 7H_2O$), is formed, which is freely soluble. The latter salt may be obtained in large rectangular prisms.* An *acid sulphate* ($C_{30}H_{24}N_2O_9, 2H_2SO_4, 7H_2O$) also exists.

The ordinary disulphate of quinia is more soluble in alcohol or alcoholic liquids than in water; hence the *Tinctura Quiniæ*, B. P., which is a solution (saturated at 55° or 60° F.) of the salt in tincture of orange-peel (eight grains in the ounce). An Ammoniated Tincture (*Tinctura Quiniæ Ammoniata*, B. P.) of similar strength is made by dissolving the sulphate in proof spirit and adding a large excess of solution of ammonia. This tincture contains quinia itself liberated from combination by the ammonia; the former tincture contains the sulphate of quinia. Quinia wine (*Vinum Quiniæ*, B. P.) is a solution of neutral sulphate and citrate of quinia in orange wine, made by dissolving the disulphate (one grain in the ounce) in orange wine by the help of citric acid. The solid official preparation of the pure disulphate is *Pilula Quiniæ*, containing three parts salt to one of confection of hips. The remaining Pharmacopœial preparation of quinia is the mixed citrates of iron, ammonium, and quinia (*Ferri et Quiniæ Citras*, B. P.), the well-known scale compound. It is made by dissolving ferric hydrate, prepared from ferric sulphate, and quinia, prepared from the sulphate, in solution of citric acid, ammonia also being added: the liquid, evaporated to a syrupy consistence and dried in thin layers on glass plates, yields the usual greenish-yellow scales (*vide* p. 170).

Basic Citrate of Quinia has the formula ($C_{30}H_{24}N_2O_9$)₃, $H_8C_6H_5O_7, 5H_2O$. Other citrates contain three molecules of quinia to two of citric acid, and one of quinia to one of citric acid.

REACTIONS.

First Analytical Reaction.—To a solution of quinia or its

* We do not know whether or not these sulphates are ordinary sulphates, the hydrogen of the acid going over to the quinia molecule, nor whether or not the quinia molecule is univalent or bivalent; hence we cannot say whether the *common sulphate* or the *soluble sulphate* is, in constitution, the neutral sulphate. In the above paragraph, the names *disulphate*, *neutral sulphate*, *acid sulphate*, indicate nothing more than that the first sulphate contains in one molecule two atoms (chemical atoms) of quinine to one of sulphuric acid, the second one of each, and the third two of acid to one of quinine.

salts in acidulated water add fresh chlorine-water, shake, and then add solution of ammonia; a green coloration (thalleioquin or *dulleiochin*) is produced. Bromine-water or bromine-vapour may be used instead of chlorine.

Second Analytical Reaction.—Repeat the foregoing reaction, but precede the addition of solution of ammonia by that of solution of ferrocyanide of potassium; an evanescent red coloration is produced (Livonius and Vogel).

Third Analytical Reaction.—To an aqueous solution of a soluble salt of quinia add solution of oxalate of ammonium; a white crystalline precipitate of oxalate of quinia falls. It is soluble in acids. If the solution to be tested be made from ordinary sulphate of quinia, excess of the latter should be added to water very faintly acidulated with sulphuric acid, and the undissolved crystals removed by filtration.

Fourth Analytical Reaction.—A saturated aqueous solution of any neutral salt of quinia is made by dissolving so much of the salt in hot water as that some shall separate when the mixture has cooled to about 60° F. After standing for some time, filter. To one volume of the filtrate one-half volume of water-washed ether is added and then ammonia in slight excess. After agitation and rest for fifteen minutes all alkaloid precipitated by the ammonia will have dissolved.

Note.—In the case of quinidia salts well defined crystals will appear at the junction of the aqueous and ethereal layers, especially after standing. In the case of cinchonidia salts a thick layer of small crystals makes its appearance at once; whilst in the case of cinchonia salts the undissolved alkaloid is enough to make the ethereal layer nearly solid.

Fifth Analytical Reaction.—Formation of Iodo-sulphate of Quinia. Dissolve sulphate of quinine in weak spirit of wine slightly acidulated with sulphuric acid and add an alcoholic solution of iodine; a black precipitate forms. Allow the precipitate to settle, pour away the fluid, wash once or twice with cold alcohol and then boil with alcohol; on cooling minute crystals separate having the optical properties of the mineral

tourmaline. This iodo-sulphate is sometimes termed Herapathite, from the name of its discoverer. It is so slightly soluble in water that by its means quinine can be separated from its admixture with the other cinchona alkaloids. According to Jörgensen it has the formula $4C_{20}H_{24}N_2O_2, 3H_2SO_4, 2HI, I_4, xH_2O$.

Sixth Analytical Reaction.—Prepare a saturated solution of ordinary sulphate of quinia in water at about 60° F., and add to 5 volumes of that solution 7 volumes of solution of ammonia (sp. gr. 0.96). The alkaloid which at first precipitates redissolves upon slight agitation if the sulphate of quinia is free from anything but traces of other cinchona alkaloids. If, however, more than traces of quinidia, cinchonidia, and cinchonina salts be present a permanent precipitate remains. This is Kerner's method of testing sulphate of quinia for other cinchona alkaloids. It turns upon the fact that the solubility of the sulphates of the cinchona alkaloids in water is in the opposite order to the solubility of the alkaloids themselves in solution of ammonia.

Other Characters.—Concentrated sulphuric acid dissolves quinia with production of only a faint yellow colour, which distinguishes it from salicin.—Quinia and its salts, heated on platinum foil, burn entirely away.—Most salts of quinia when in solution have a beautiful blue fluorescence. They twist the ray of polarized light to the left. Quinia is soluble in alcohol, ether, benzol and chloroform. Ordinary quinia sulphate is insoluble in chloroform, and but slightly soluble in water. Its solubility in chloroform is increased by the presence in solution of quinidia and cinchonina sulphates (Prescott), and its solubility in water is decreased by the presence in solution of ammonium sulphate (Carles). The slight solubility of its sulphate and iodo-sulphate in water distinguishes quinia from the other cinchona alkaloids, including the "amorphous alkaloid," or "quinoidia."

Quinidia or Quinidine ($C_{20}H_{24}N_2O_2$, the Conquinine or Con-

chinine of Hesse), is, an isomer of quinia. Its salts are fluorescent, and give thalleioquin with chlorine- or bromine-water and ammonia. They twist the ray of polarized light to the right. Quinidia is insoluble in water and sparingly soluble in ether (*see* Quinia, 4th Analytical Reaction). It is soluble in alcohol, benzol, and chloroform. It is less soluble than quinia in ammonia, 5 volumes of a saturated aqueous solution of its ordinary sulphate requiring 60 to 80 volumes of ammonia solution (sp. gr. 0.96). Its sulphate is more soluble in water and chloroform than the sulphate of quinia. Tartrate of quinidia is soluble in water. The hydriodate is insoluble in water and weak spirit, and occurs as sandy crystals. The hydriodates of the other cinchona alkaloids, though more soluble than quinidin hydriodate, are sometimes precipitated from neutral concentrated solutions, as amorphous or semiliquid precipitates. These, however, are soluble in weak spirit.

Cinchonidia or *Cinchonidine* ($C_{20}H_{24}N_2O$).—When perfectly pure, salts of cinchonidia do not give thalleioquin and are not fluorescent. Even good commercial salts, however, nearly always give both reactions. Salts of cinchonidia twist the polarized ray to the left. Cinchonidia is insoluble in water and nearly so in ether (*see* Quinia, 4th Analytical Reaction). It is soluble in alcohol, benzol, and chloroform. It is less soluble in ammonia solution than quinia, five volumes of a saturated aqueous solution of cinchonidia sulphate requiring about 80 volumes of ammonia solution (sp. gr. 0.96). Sulphate and hydriodate of cinchonidia are soluble in water, but the sulphate, like quinia sulphate, is insoluble in chloroform. Tartrate of cinchonidia is insoluble in water, and in this form cinchonidia is usually separated from neutral solutions containing the other cinchona alkaloids except quinia.

Cinchonia or *Cinchonine* ($C_{20}H_{24}N_2O$) is an isomer of cinchonidia. When quite pure its salts are not fluorescent and do not give thalleioquin. As in the case of cinchonidia even good commercial specimens of cinchonia salts nearly always give both reactions. Cinchonia salts twist the polarized ray to the right. Cinchonia is insoluble in water and nearly so in ether (*see* Quinia, 4th Analytical Reaction). It is soluble in chloroform, benzol, and alcohol. Chloroform containing one-fourth of its weight of 95% alcohol dissolves cinchonia much more readily than either alcohol or chloroform alone. Cinchonia is insoluble in ammonia solution. Sulphate, tartrate,

and hydriodate of cinchonia are soluble in water, and the sulphate, like sulphate of quinidia, is soluble in chloroform. In mixtures of cinchona alkaloids, this alkaloid is precipitated by alkali after the others have been successively removed by ether, tartrate of sodium and iodide of potassium.

"*Quinoidia*" or "*Quinoidine*," or the "*Amorphous Alkaloid*."—Cinchona barks generally contain some alkaloid isomeric with quinia which like quinia is soluble in ether, but the ordinary sulphate and iodosulphate of which are not crystalline and are soluble. These salts are semisolid resinous looking substances. The iodo-sulphate is used in De Vrij's method for the separation of mixed alkaloids (*see Index*). Quinoidia is usually obtained along with quinia, etc., from the mixed alkaloids by ether, and remains in the mother-liquor, from which it is precipitated by an alkali.

Cinchovatia or *Cinchovatine* occurs in a particular variety of cinchona-bark. *Quinicia* or *Quinicine* and *Cinchonicia* or *Cinchonicine* are alkaloids produced by the action of heat on quinia or quinidia and on cinchonia or cinchonidia respectively. They, also, are isomers, Hesse says polymers, of the parent alkaloids. Both yield ordinary salts. *Quiniretin* is the name given to the brown or reddish-brown indifferent substance into which quinine in aqueous solution is converted when exposed to much light.

Quinamia or *Quinamine* ($C_{20}H_{26}N_2O_2$) is a fifth cinchona alkaloid obtained by Hesse in 1872 from the bark of *Cinchona succirubra*. Its solution is not fluorescent and does not give thalleioquin.

STRYCHNIA, OR STRYCHNINE.

Formula $C_{31}H_{22}N_2O_2$. Molecular weight 334.

Source.—This alkaloid exists, to the extent of 0.2 to 0.5 per cent., in *Nux Vomica* (*Strychnos Nux Vomica*), also (Shenstone) in minute quantity in the bark of the *Nux Vomica* tree (false angostura bark), and to 1.0 or 1.5 per cent. in *St. Ignatius's bean* (*Strychnos Ignatius*), chiefly in combination with strychnic or igasuric acid, or, after slight fermentation when moistened, with lactic acid.

Process.—According to the official process for its preparation (*Strychnia*, B. P.), the nuts, disintegrated by subjection to steam, and, after drying, grinding in a coffee-mill, are ex-

hausted with spirit, the latter removed by distillation, the extract dissolved in water, colouring and acid matters precipitated by acetate of lead, the filtered liquid evaporated to a small bulk, the strychnia precipitated by ammonia, the precipitate washed, dried, and exhausted with spirit, the spirit recovered by distillation, and the residual liquid set aside to crystallize. Crystals of strychnia having formed, the mother-liquor (which contains the brucia of the seeds) is poured away, and the crystals of strychnia washed with spirit (to remove any brucia) and recrystallized.

Properties.—Strychnia occurs "in right square octahedrons or prisms, colourless and inodorous; sparingly soluble in water, but communicating to it its intensely bitter taste; soluble in boiling rectified spirit and in chloroform, but not in absolute alcohol or in ether."

REACTIONS.

First Analytical Reaction.—Place a minute particle of strychnia on a white plate, and near to it a small fragment of red chromate of potassium; to each add one drop of concentrated sulphuric acid: after waiting a minute or so for the chromate to fairly tinge the acid, draw the latter, by a glass rod, over the strychnia spot; a beautiful purple colour is produced, quickly fading into a yellowish red. The following oxidizing agents may be used in the place of the chromate:—puce-coloured oxide of lead, fragments of black oxide of manganese, ferridcyanide of potassium, or permanganate of potassium.

This reaction is highly characteristic; a minute fragment dissolved in much dilute alcohol, or, better, chloroform, and one drop of the liquid evaporated to dryness on a porcelain crucible-lid or other white surface, yields a residue which immediately gives the purple colour on being oxidized in the manner directed.

Other Reactions.—Strong sulphuric acid does not act on strychnia, even at the temperature of boiling water, a fact of which advantage is taken in separating strychnia from other organic matter for the purposes of toxicological analysis.—Sulphocyanide of potassium produces, even in dilute solutions of strychnia, a white precipitate, which, under the

microscope, is seen to consist of tufts of acicular crystals.—Strong nitric acid does not colour strychnia in the cold, and on heating only turns it yellow.

The Physiological Test.—A small frog placed in an ounce of water to which $\frac{1}{1000}$ of a grain of a salt (acetate) of strychnia is added, is, in two or three hours, seized with tetanic spasms on the slightest touch, and dies shortly afterwards.

Strychnia has an intensely bitter taste. Cold water dissolves only $\frac{1}{10000}$ part; yet this solution, even when largely diluted, is distinctly bitter. Alcohol is a somewhat better solvent. The salts of the alkaloid are more soluble. The official solution (*Liquor Strychniæ*, B. P.) contains four grains of strychnia to the ounce, the solvent being three parts water, one part spirit, and a few minims (6 per ounce) of hydrochloric acid (rather more than sufficient to form hydrochlorate of strychnia).

BRUCIA, or BRUCINE ($C_{23}H_{26}N_2O_4, 4H_2O$), is an alkaloid accompanying strychnia in *Nux Vomica* and St. Ignatius's bean to the extent of about 0.5 per cent. It is readily distinguished by the intense red colour produced when nitric acid is added to it. *Igasuria*, once supposed to be a third alkaloid of *nux vomica*, has been shown by Shenstone to be only a mixture of brucia and strychnia.

Curarine ($C_{10}H_{15}N$), the active principle of the arrow-poison termed *curari*, *urari*, *ourari*, *wourali*, or *woorara*, prepared from a *strychnos*, resembles strychnia in giving colour by oxidation, but the colour is more stable. Iodide or platino-cyanide of potassium do not with curarine afford precipitates which crystallize from alcohol like those of strychnia. Curarine, also, is soluble in water. Unlike strychnia curarine is reddened by sulphuric acid; it, also, is not dissolved out by ether from an acid or alkaline liquid. *Ourari* appears to vary much in strength and quality. It is probably a mixture of vegetable extracts.

Distinction of Brucia from Morphia.—The red coloration produced by the action of nitric acid on brucia is distinguished from that yielded by morphia by the action of reducing agents (such as stannous chloride, hyposulphite of sodium, sulphhydrate of sodium), which decolourize the morphia-red, but change that of the brucia to violet and green (Cotton).

Distinction of free alkaloids or their salts from each other.—This is accomplished by remembering the appearance and other physical characters of the substances as met with in phar-

macy, the effect of heat, the action of such solvents as water, alcohol, and ether, the influence of strong and diluted acids, strong and weak alkalies, oxidizing substances, and other reagents. (See annexed Tables, 1 and 2.)

QUESTIONS AND EXERCISES.

678. What alkaloids are more or less characteristic of the different varieties of cinchona-bark? In what form do they occur?

679. By what method is Disulphate of Quinia obtained?

680. Give the characters of disulphate of quinia.

681. Describe the tests for quinia.

682. How would the adulteration of disulphate of quinia by salicin be detected?

683. Show how the sulphates of quinidia or cinchonina may be proved to be present in commercial quinia.

684. How are cinchonina and quinia distinguished from morphia?

685. Whence is Strychnia obtained?

686. Describe the official process for the isolation of strychnia.

687. Give the characters of strychnia.

688. Enumerate the tests for strychnia, and describe their mode of application.

689. By what reagent is brucia distinguished from strychnia?

690. Distinguish between brucia and morphia.

691. By what general methods would you distinguish common alkaloids from each other?

Analytical Exercises.—Analyse small quantities of alkaloids, their salts, and various "scale" compounds by aid of the annexed Tables, 1 and 2.

ALKALOIDS OF LESS FREQUENT OCCURRENCE.

ACONITIA, ACONITINA, or ACONITINE is an alkaloid obtained from aconite (*Aconitum Napellus*) leaves (*Aconiti Folia*, B. P.) and root (*Aconiti Radix*, B. P.). The alkaloid itself is only slightly soluble in water; it occurs in the plant in combination with a vegetable acid, forming a soluble salt.

Process.—The official process for its preparation (*Aconitia*,

IDS, ACIDS, ETC.

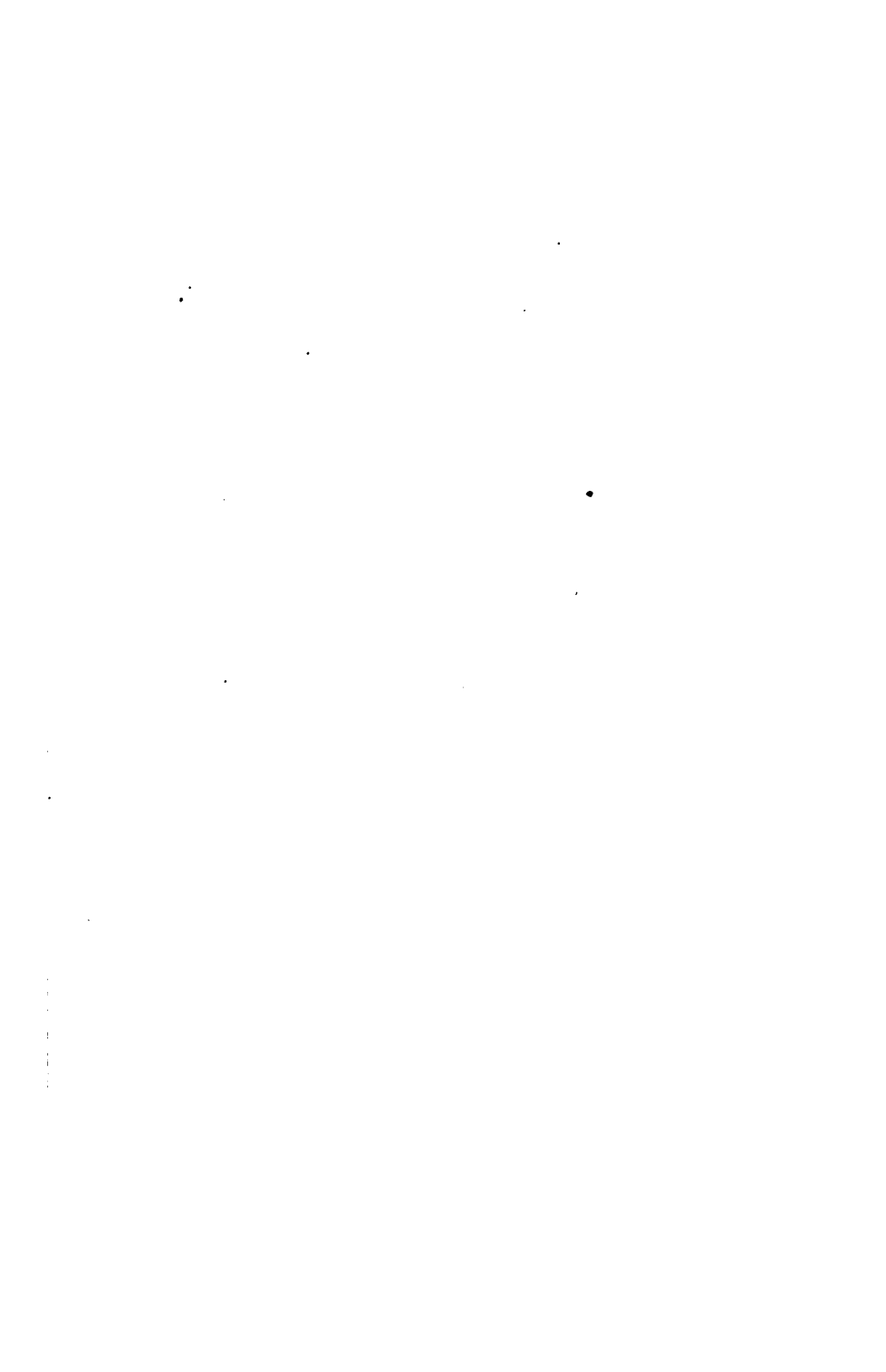
MORPH. BRUC. SULPHURIC. HYDROCHLORIC. PHOSPHORIC. ACETIC. MECONIC. CITRIC.					
To a small quawith a slight excess of white plate add stinto three portions, and acid.		Make a neutral solution, divide it into two portions, and test as follows :—			
<i>An orange colour, de-colourised by SnCl_2, $\text{Na}_2\text{S}_2\text{O}_3$, or NaHS = <i>Morphia</i>.* Confirm by—Fe_2Cl_6 (neut.), which gives with morphia or its salts a blue colour. HIO_3 is decomposed by morphia or its compounds, with liberation of iodine which may be recognised by starch.</i>		<i>And portion.</i> grs AgNO_3, tbl precipi- dro = <i>Hydro- Hic Acid</i>. to dil was ad dro not A ou	<i>Third por- tion.</i> Boil for a time and then add a few drops to an excess of solution of molybdate of ammonium in HNO_3 and boil. A yellow pulverulent precipitate = <i>Phosphoric Acid</i>.	<i>First por- tion.</i> Add neutral Fe_2Cl_6. A red colour indicates acetic or meconic acid. Distinguish Decolourised by boiling = <i>Acetic Acid</i>. Not decolourized by boiling = <i>Meconic Acid</i>.	<i>Second por- tion.</i> If no acetic or meconic acid, add AgNO_3. Precipitate (white, with a tendency to darken) soluble in HNO_3 = <i>Citric Acid</i>. Confirm by adding CaCl_2 to a neutral conc. solution and boiling, when a white precipitate of citrate of calcium separates.
* Strychnia of commerce often gives an orange or red colour, due to contamination with brucia.					

[Table 1. To

LE COMPOUNDS.

	pyrophosphoric).	INORGANIC BASES. (AMMONIUM (often as a contamination). FERRIC IRON. POTASSIUM. SODIUM (rare).
Dissolve the ash with HNO_3 , with a little polybdate in HNO_3 aqueous solution		<i>Ammonium</i> —Boil aqueous solution of scale with KHO and test vapour for NH_3 . Filter and dissolve precipitate in HCl and test the solution for Iron by K_4FeCy_6 , KCyS , etc. <i>Potassium and Sodium</i> .—Ignite a small quantity of the scale, and moisten the residue with water. Test moistened residue with litmus paper. If alkaline examine for potassium and sodium by the colour imparted to flame, and for potassium by HCl and PtCl_4 .
ETHER PRECIPITATE.		
May contain of the aqueous beberia. To H_2O , filter, and add H_2O , the filtrate a slight with HC_2H_3 and divide into two the ether. To BaCl_2 (precipitate solution, add). To the other add $\text{NH}_4\text{Bate} = \text{Hydrochloric}$ another portion of with HNO_3 and add		
GREEN CO (thalleog)	PRECIPITATE WHITE.	
Solution in cent, and either quinia dia. Concent remainder of tion and div two parts. add KI , and other add (NH_4I precipitates quinia. For another see the third of the annex (1) for alkaloi 40 or 50 g material.	<i>Citric acid</i> gives no mirror. CaCl_2 and Ca_2HO do not precipitate citric acid in the cold, but upon boiling (if solution be sufficiently concentrated) precipitation occurs.	
	<i>Citric Acid</i> .—To slightly acidified KHO filtrate, slight excess and considerable quantity of NH_4Cl filtrates are precipitated completely in the cold with for about ten minutes. To the solution (or filtrate, present) add three volumes of spirits of wine when precipitated. If sulphates have been found disregard with spirits of wine.	

[Table 2.]



B. P.) consists in dissolving out the natural salt of the alkaloid from the root by rectified spirit, recovering the latter by distillation, mixing the residue with water, filtering, precipitating the aconitia by ammonia, drying the precipitate and digesting it in ether (in which some of the accompanying impurities are insoluble), recovering the ether by distillation, dissolving the dry residue in the retort in water acidulated by sulphuric acid, again precipitating the alkaloid by ammonia, and finally washing and drying.

Properties.—Aconitia usually occurs as a white powder, but has been obtained and studied in the crystalline state by Groves, Wright, and others. It is soluble in 150 parts of cold water, 50 of hot, and much more soluble in alcohol and in ether. It is one of the most violent poisons known. "When rubbed on the skin it causes a tingling sensation, followed by prolonged numbness." The thousandth part of a grain on the tip of the tongue produces, after a minute or so, a characteristic tingling sensation and numbness; large quantities rubbed into the skin cause numbness and loss of feeling. Oil of vitriol turns it of a yellowish and, afterwards, dirty violet colour.

According to Wright, who, in conjunction with Groves and Williams, worked by the aid of grants from the British Pharmaceutical Conference, *Aconitum napellus* yields, chiefly, crystalline aconitine, $C_{33}H_{43}NO_{13}$, with some crystalline pseudaconitine, $C_{36}H_{49}NO_{13}$ and a little non-crystalline alkaloid.

The tuberous roots of *Aconitum ferox* and other species constitutes the *bish* or *bikh* of India (*Aconiti ferocis Radix*, P. I.). It chiefly contains the variety of aconitine termed *pseudaconitine*. Some of the aconitine of pharmacy is pseudaconitine.

According to Paul and Kingzett the alkaloid of Japanese aconite has the formula $C_{29}H_{43}NO_6$, while Wright and Menke state that the formula is $C_{36}H_{38}N_2O_{31}$, and name it *japaconitine*.

Unguentum Aconitiæ, B. P., contains eight grains of the alkaloid to one ounce of prepared lard.

Aconitum heterophyllum, *Atis*, or *Atees* (*Aconiti heterophylli Radix*, P. I.) contains no aconitia, but an alkaloid having the formula $C_{46}H_{74}N_2O_5$ (Broughton, in "Pharmacographia").

ATROPIA, or ATROPINE ($C_{17}H_{23}NO_3$), exists in the *Belladonna*, or Deadly Nightshade (*Atropa Belladonna*; *Belladonnæ Folia et Radix*, B. P.), as soluble acid malate of atropia.

Process.—It is obtained in the pure state by exhausting the

root with spirit, precipitating the acid and some colouring-matter by lime, filtering, adding sulphuric acid to form sulphate of atropia (which is somewhat less liable to decomposition during subsequent operations than the alkaloid itself), recovering most of the spirit by distillation, adding water to the residue, and evaporating till the remaining spirit is removed; solution of carbonate of potassium is then poured in till the liquid is nearly, but not quite, neutral, by which resinous matter is precipitated; the latter is filtered away, excess of carbonate of potassium then added, and the liberated atropia dissolved out by shaking the liquid with chloroform. The latter solution, having subsided, is removed, the chloroform recovered by distillation, the residual atropia dissolved in warm spirit, colouring-matter separated by digesting the liquid with animal charcoal, the solution filtered, evaporated, and set aside to deposit crystals.

Solubility.—Atropia is sparingly soluble in water, the liquid giving an alkaline reaction—more soluble in alcohol and ether.

Tests.—Atropia solutions give with perchloride of gold a yellow precipitate. One drop of a dilute aqueous solution (two grains to the ounce) powerfully dilates the pupil of the eye. It is generally applied on a piece of thin tissue paper or small disk placed between the eyelid and the eye. Baryta water decomposes it into *tropine* ($C_8H_{15}NO$), and *tropic acid* ($C_9H_{10}O_3$) a molecule of water being absorbed, hence the atropia, so-called, would seem to be *tropate of tropine*. Indeed Ladenburg by combining tropic acid and tropine has produced a base indistinguishable from atropia. The same chemist by removing the elements of water from tropine gets *tropidine*, $C_8H_{13}N$. This is, possibly, an intermediate member of a group of monamines of which others are *conine*, $C_8H_{15}N$, and *collidine* $C_8H_{11}N$, the latter a product of the destructive distillation of bone-oil, coal, quinine, etc.

Preparation.—The alkaloid itself (*Atropia*), its sulphate (*Atropiæ Sulphas*, a colourless powder soluble in water, made by neutralizing atropia with sulphuric acid), their solutions (*Liquor Atropiæ*, four grains per ounce, and *Liquor Atropiæ Sulphatis*, four grains per ounce), and an ointment (*Unguentum Atropiæ*, eight grains per ounce) are the preparations official in the British Pharmacopœia.

BEBERIA, BEBERINE, BEBIRINE, or BIBIRINE ($C_{18}H_{27}NO_3$), is an alkaloid existing in the bark of Bebeeru, or Bibiru (*Nectandra Rodiæi*).

Process.—According to the British Pharmacopœia, it, or rather its sulphate, $C_{36}H_{49}N_3O_6$, H_2SO_4 (*Beberia Sulphas*, B. P.), may be prepared by exhausting the bark (*Nectandra Cortex*, B. P.) with water acidulated by sulphuric acid, concentrating, removing most of the acid by lime, filtering, precipitating the alkaloid by ammonia, filtering, drying, dissolving in spirit (in which some accompanying matters are insoluble), recovering most of the spirit by distillation, neutralizing by dilute sulphuric acid, evaporating to dryness, dissolving the residual sulphate in water, evaporating to the consistence of a syrup, and spreading on glass plates, drying the product at $140^\circ F$. Thus obtained, it occurs in thin dark-brown translucent scales, yellow when powdered, strongly bitter, soluble in water and in alcohol.

Tests.—Alkalies give a pale yellow precipitate of beberia when added to an aqueous solution of a salt of the alkaloid; the precipitate is soluble in ether. With red chromate of potassium and sulphuric acid beberia gives a black resin, and with nitric acid a yellow resin.

Buzine, from the bark of *Buxus sempervirens*; *Pelosine*, or *Cissampeline*, from the root (*Pareira Radix*, B. P.) of *Cissampelos Pareira*; and *Paricine*, from a false Para cinchona-bark, are probably identical with Beberine (Flückiger).

Nectandria ($C_{20}H_{23}NO_4$).—Drs. MacLagan and Gamgee a few years ago discovered this second alkaloid in Bebeeru-bark. It differs from beberia in fusing when placed in boiling water, in being much less soluble in ether, in giving with strong sulphuric acid and black oxide of manganese a beautiful green and then violet coloration, and in having a distinct molecular weight. They gave the opinion that two other alkaloids exist in Bebeeru-bark.

BERBERIA, or BERBERINE ($C_{20}H_{17}NO_4$), is an alkaloid existing in several plants of the natural order *Berberideæ* (three species yield *Indian Barberry*, *Berberis Cortex*, P. I.), in Calumba-root (*Calumbæ Radix*, B. P.), in the root of *Coptis Teeta*, or *Mishmi Bitter* (*Coptidis Radix*, P. I.), an Indian tonic, and in many other yellow woods. *Hydrastis canadensis*, *Yellow root*, or *Golden seal*, contains berberia, though another alkaloid, *hydrastia*, and even a third, are said by Hale to be present. *Xanthorrhiza apiifolia*, an old American tonic, and, apparently, *Xanthoxylon fraxineum*, or *Prickly Ash*, also contain berberia. The rhizome of *Menispermum Canadense*, *Yellow Parilla*, contains, according to Maisch, a colourless alkaloid

as well as berberia. The colour of the tissues of these vegetables is apparently due to berberia; for the alkaloid itself is remarkable for its beautiful yellow colour.

Tests.—When a dilute solution of iodine in iodide of potassium is added to a solution of any salt of berberia in hot spirit, excess of iodine being carefully avoided, brilliant green spangles are deposited. The reaction is sufficiently delicate to form, according to Perrins, an excellent test of the presence of berberia. This iodo-compound polarizes light, and has other analogies with a similar quinine-salt termed herapathite.

Berberia is not an official alkaloid; but the plants in which it occurs are used as medicinal agents in all parts of the world.

Process.—Berberia is readily extracted by boiling the raw material with water, evaporating the strained liquid to a soft extract, digesting the residue in alcohol, recovering the alcohol by distillation, boiling the residue with diluted sulphuric acid, filtering and setting aside; the sulphate of berberia separates out, and may be purified by recrystallization from hot water. The alkaloid itself is obtained by shaking hydrate of lead with a hot aqueous solution of the sulphate of berberia (Procter).

CAPSICUM, or CAPSICINE.—M. Felletár obtained from capsicum-fruits (*Capsici Fructus*, B. P.) a volatile alkaloid having the smell of conia. Thresh has obtained crystalline hydrochlorate and sulphate. The latter chemist has also succeeded in isolating the active principle of capsicum, which he has termed *capsaicin* ($C_9H_{14}O_2$), a crystalline non-alkaloidal excessively acrid substance. Its exact chemical character is not yet made out. (See also *Capsicin*, in Index.)

COCAINA, or COCAINE ($C_{18}H_{19}NO_4$) is an alkaloid of *Erythroxylon Coca*, the leaves of which are said to be a powerful restorative to the human system.

COLCHICUM, or COLCHICINE, the active principle of *Colchicum autumnale* (*Colchici Cormus*; *Colchici Semina*, B. P.), is said to be an alkaloid, though some investigators think it has more of the characters of a neutral substance and give it the name *Colchicin*. It needs further examination. The most active medicinal preparation is an extract made from the fresh seeds by digestion in large volumes of alcohol of at least 90 per cent., and subsequent digestion of the marc in hot water. The extracts left on evaporating the two fluids separately are to be carefully mixed (Mols).

CONIA, CONYLIA, CONINE, CONICINE, or CICUTINE.—Formula $C_8H_{15}N$, or $(C_8H_{14})^+HN$. This alkaloid is a volatile liquid, occurring in hemlock (*Conium maculatum*) in combination with an acid (malic?). It is not official. According to Petit its boiling point is $170^\circ C.$, and its density 0.846.

Process.—It may be obtained by distilling hemlock-fruit (*Conii Fructus*, B. P.) with water rendered slightly alkaline by caustic soda or potash, or by similarly treating the fresh juice of the leaves. The alkaloid is a yellow oily liquid, floating on the water that distils over; by redistillation it is obtained colourless and transparent.

The salts of conia have no odour, but when moistened with solution of an alkali yield the alkaloid, the strong smell of which, at once recalling hemlock, is characteristic. Extract of hemlock-leaves (*Conii Folia*, B. P.), to which solution of potash and boiling water have been added, forms the official Inhalation of Conia (*Vapor Conice*, B. P.).

Tests.—Sulphuric acid turns conia purplish red, changing to olive-green, nitric acid a blood-red; perchloride of gold produces a yellowish-white precipitate, perchloride of platinum no precipitate, in aqueous solutions.

Hemlock also contains *methyl-conia* $(C_8H_{14})^+CH_3N$ (Kekulé and Von Planta) and *conhydrine*, $NC_8H_{14}H_2O$.

According to Schiff, conia, isomeric, at least, with the natural alkaloid, may be produced artificially by action of ammonia on butyric aldehyd and destructive distillation of the resulting compound.

CORYDALIA, or CORYDALINA, or CORYDALINE is an alkaloid obtained by Wenzell from "Turkey Corn," the tubers of *Dicentra (corydalis) formosa*.

DATURIA, or DATURINE, *vide* HYOSCYAMIA.

DELPHIA, DELPHINE, or DELPHININE $(C_{24}H_{35}NO_3)$, the poisonous alkaloid of Stavesacre (*Delphinium staphysagria*). The powdered seeds of the plant are employed to kill the *pediculi* of animals.

DITAMINE (Jobst and Hesse), present in the *Ditain* of Gruppe, is an alkaloid of "Dita," or bark of *Echites scholaris*, or *Alstonia scholaris* (*Alstonia Cortex*, P. I.), a reputed febrifuge. Others are *echitamine* and *echitenine*. Oberlin and Schlagdenhauffen state that the allied *Alstonia constricta* contains a crystalline alkaloid *alstonine*, and uncrystallizable *alstonicine*.

EMETIA, or EMETINE $(C_{30}H_{44}N_2O_4)$; Glenard).—This alkaloid is the active emetic principle of *Oephalis ipecacuanha* (*Ipecacu-*

anha, B. P.). It occurs in combination with ipecacuanhic acid. The nitrate is peculiarly slightly soluble in water (Lefort). In the *Pulvis Ipecacuanhæ Compositus*, B. P. and U. S. P., or "Dover's Powder" (Powdered Ipecacuanha, 1 part; Powdered Opium, 1 part; and Sulphate of Potassium, 8 parts), minute division of the active ingredients is promoted by prolonged trituration with sulphate of potassium, which is a very hard salt.

The Indian substitute of Ipecacuanha is the dried leaf (*Tylophora Folia*, P. I.) of *Tylophora asthmatica*. Its active principle has not been satisfactorily isolated.

GELSEMINIA, or GELSEMINE ($C_{11}H_{19}NO_3$) is the alkaloid of *Gelsemium sempervirens*, or Carolina yellow jasmine (*Gelsemium*, U. S. P.) in the tissues of which plant *æsculin* ($C_{30}H_{34}O_{19}$, the *gelseminic acid* of Wormley), the fluorescent glucoside of the Horse Chestnut, and of many other plants, is also present.

HYOSCYAMIA, or HYOSCYAMINE ($C_{17}H_{23}NO_3$), a volatile alkaloid occurring in the leaves (*Hyoscyami Folia*, B. P.) and other parts of Henbane. Its effect on the eye is similar to that of atropia. The researches of Buckheim point to the conclusion that hyoscyamine is the tropate of an alkaloid homologous with tropine (Sonnenschein). (See ATROPIA.)

The alkaloids which occur in *Datura Stramonium*, or Thorn apple (*Stramonii Folia et Semina*, B. P.), *Dhatura* (*Datura alba*; *Datura Folia et Semina*, P. I.), and in *Duboisia Myoporoides*, and were formerly supposed to be distinct alkaloids, called respectively *Daturia* and *Duboisia*, are identical with Hyoscyamia, and the latter is isomeric if not identical with atropia (Ladenburg). Bees which sip from the flowers of stramonium are said to produce poisonous honey.

JERVIA, or JERVINE ($C_{30}H_{46}N_2O_8$), occurs in *Veratrum album*, White Hellebore, and *V. viride**, American White Hellebore, the root of which is officially recognised in Great Britain (*Veratri viridis Radix*, B. P.). Its salts are much less soluble in water than those of veratria. According to Bullock, *Veratrum viride* contains still another alkaloid—*veratroidia*; and, according to Mitchell, *Veratrum album* also contains an alkaloid which he terms *veratralbia*. Tobien gives the formula of jervia as $C_{27}H_{47}N_2O_8$, and of veratroidia as $C_{51}H_{78}N_2O_{16}$, or

* "The name *Green Hellebore* is sometimes applied to the drug; but it properly belongs to *Helleborus viridis* (p. 497), which is medicinal in some parts of Europe."—Hanbury.

$C_{24}H_{37}NO_7$. According to Wright, *Veratrum album* contains jervine, $C_{28}H_{37}NO_8$; pseudojervine, $C_{29}H_{43}NO_7$; rubijervine, $C_{26}H_{43}NO_8$; veratralbine, $C_{28}H_{43}NO_6$; and traces of veratrine, $C_{37}H_{63}NO_{11}$. The same author finds *Veratrum viride* to contain jervine, pseudojervine, cevadine $C_{32}H_{49}NO_9$; rubijervine and traces of veratrine and veratralbine.

LOBELINA, or LOBELINE.—A volatile fluid alkaloid first isolated from the dried flowering herb *Lobelia inflata* (*Lobelia*, B. P.) by Procter. In the pure state it smells slightly of the plant, but mixed with ammonia it emits a strong and characteristic smell of the herb. With acids it forms crystalline salts.

LUPULINE is stated by Greismayer to be a liquid volatile alkaloid contained in Hops (*Humulus lupulus*).

NECTANDRIA (*vide* BEBERIA).

NICOTIA, NICOTINA, NICOTYLIA, or NICOTINE.—Formula $C_{10}H_{14}N_2$, or $(C_5H_7)_2N_2$. This is also a volatile liquid alkaloid, forming the active principle of tobacco (*Nicotiana tabacum*), malate and citrate of nicotina being the forms in which it occurs in the leaf (*Tabaci Folia*, B. P.). Its odour is characteristic; like conia, it yields a precipitate with perchloride of gold; but, unlike that alkaloid, its aqueous solutions are precipitated yellowish white by perchloride of platinum. It is not official. It is also contained in *Pituri*, a drug "chewed by the natives of some parts of Australia as a stimulant narcotic."

PHYSOSTIGMIA, or PHYSOSTIGMINE ($C_{30}H_{31}N_3O_4$). An alkaloid contained in the Calabar Bean (*Physostigmatis Faba*), the seed of *Physostigma venenosum* (Jobst and Hesse). A trace of it powerfully contracts the pupil of the eye; a small quantity is highly poisonous. Fraser also isolated another (but, possibly, the same) principle, and termed it *Eseria*, from Eseré, the name of this ordeal-poison at Calabar.

PILOCARPINE is, apparently, the active principle of the diaphoretic and sialogogue *Jaborandi* (*Pilocarpus pennatifolius*). The occurrence of an alkaloid in this plant was first announced by Hardy, followed almost immediately by Byasson. A crystalline nitrate and hydrochlorate were first obtained by Gerrard. Kingzett gives the formula as $C_{23}H_{34}N_4O_4$. The leaves also yield an essential oil, a terpene, $C_{10}H_{16}$ (Hardy).

PIPERIA, or PIPERINE ($C_{17}H_{19}NO_3$), is a feeble alkaloid occurring in white, black (*Piper Nigrum*, B. P.), long pepper (*Chanica officinarum*, Mign), and cubeb pepper (*Cubeba*, B. P.) asso-

ciated with volatile oil and resin; to these three substances the odour, flavour, and acidity belong. Piperia is obtained on boiling white pepper with alcohol, and evaporating the liquid with solution of potash, which retains resin. Recrystallized from alcohol, piperia forms colourless prisms fusible at 212° F. With acids and certain metallic compounds it forms salts, and distilled with strong alkali yields *piperidia* or *piperidine* ($C_5H_{10}HN$), an alkaloid of strong chemical properties, and *piperic acid* ($C_{12}H_{10}O_4$). According to Buckheim the amorphous resin of the peppers is similar in constitution to piperine, alkalies breaking it up into piperidine and *chavicol acid*. *Pyrethrin* is also said to be a member of the series. The piperine of cubeb pepper is not to be confounded with *cubebin*, a neutral constituent having the formula $C_{33}H_{34}O_{10}$.

SANGUINARINA is the alkaloid of *Blood Root* (*Sanguinaria canadensis*). Its salts are red.

SOLANIA, or SOLANINE ($C_{45}H_{69}NO_{16}$).—An alkaloid said to exist in the Woody Nightshade or Bitter-sweet (*Solanum dulcamara*). The dried young branches of the plant are official (*Dulcamara*, B. P.). This alkaloid is only slightly soluble in water, alcohol, or ether; nitric acid colours it yellow; sulphuric acid produces at first a yellow, then a violet, and finally a brown coloration. It is said to be a conjugated compound of sugar with *solanidine* ($C_{25}H_{39}NO$). Geissler finds *dulcamarin* ($C_{33}H_{34}O_{10}$), a glucoside, to be the bitter constituent of *Solanum dulcamara*.

SPARTEIA, or SPARTEINE ($C_{15}H_{26}N$), is a poisonous volatile alkaloid occurring in Broom-tops (*Scoparii Cacumina*, B. P.). Its discoverer, Stenhouse, considers that the diuretic principle of broom is *Scoparin*, a non-poisonous body. Mills has obtained ethyl-sparteine ($C_{15}H_{25}C_2H_5N$) and diethyl-sparteine ($C_{15}H_{24}C_2H_5C_2H_5N$).

THEIA, THEINE, or CAFFEINE ($C_8H_{10}N_4O_2 + H_2O$).—This alkaloid occurs in tea, 1 to 4 %; coffee, .5 to 2 %; Maté or Paraguay tea, .2 to 2 %; guarana, 5 %; and the kola-nut. Infusions and preparations of these vegetable products are used, chiefly as beverages, by three-fourths of the human race. It is remarkable that the instinct of man, even in his savage state, should have led him to select, as the bases of common beverages, just the four or five plants which out of many thousands are the only ones, so far as we know, containing theia.

Theia is volatile. Considerable quantities may be collected

by condensing the vapours evolved during the roasting of coffee on the large scale. The infusion of tea, from which astringent and colouring-matters have been precipitated by solution of subacetate of lead, and which has been evaporated to a small bulk, yields a precipitate of theine on the addition of a strong solution of carbonate of potassium. It may be crystallized from alcohol or by sublimation.

Test.—Concentrated nitric acid, or a mixture of chlorate of potassium and hydrochloric acid, rapidly oxidizes theia, forming compounds which with ammonia yield a beautiful purple-red colour, resembling the murexid obtained under similar circumstances from uric acid; the oxidation must not be carried too far. Theine boiled with caustic potash yields methylamine (CH_3HHN), the vapour of which has a peculiar, characteristic odour.

The chemical action of theia on the system is not yet made out. It is probably a pure stimulant.

VERATRIA, or VERATRINE ($\text{C}_{32}\text{H}_{50}\text{NO}_9$, Schmidt and Köppen; $\text{C}_{53}\text{H}_{86}\text{N}_2\text{O}_{15}$, Weigelin).—This alkaloid occurs in *Cevadilla* (*Sabadilla*, B. P.; the seeds of *Asagrea officinalis*, Lindley, termed *Sabadilla officinarum* by Brandt, and *Veratrum officinale* by Schlecht). It is also said to occur in the leaves of *Sarracenia purpurea*. According to Weigelin, *Cevadilla* contains two isomeric varieties of veratria, the one soluble the other insoluble in water. He says there are also present *Sabadilline* ($\text{C}_{41}\text{H}_{66}\text{N}_2\text{O}_{13}$) and *Sabatrine* ($\text{C}_{51}\text{H}_{86}\text{N}_2\text{O}_{17}$). The veratrine of trade contains the two latter alkaloids (Weigelin). A mere trace of veratria brought into contact with the mucous membrane of the nose causes violent fits of sneezing. These alkaloids and those from the different species of veratrum are evidently very closely allied. Wright and Luff, by the use of tartaric acid, a solvent less likely than the stronger acids to decompose alkaloids, extract from *Cevadilla*, *Veratrine* ($\text{C}_{37}\text{H}_{53}\text{NO}_{11}$), *Cevadine*, $\text{C}_{32}\text{H}_{49}\text{NO}_9$; and *Cevadilline*, $\text{C}_{34}\text{H}_{53}\text{NO}_8$.

The official process for the preparation of the alkaloid (*Veratria*, B. P.) consists in exhausting the disintegrated *cevadilla*-seeds by alcohol, recovering most of the spirit by distillation, pouring the residue into water, by which much resin is precipitated, filtering, and precipitating the veratria from the aqueous solution by ammonia. It is purified by washing with water, solution in dilute hydrochloric acid, decolorization of the liquid by animal charcoal, reprecipitation by ammonia, washing and drying.

Unguentum Veratriæ, B. P., contains eight grains of the slightly impure alkaloid obtained as just described, rubbed down with half a drachm of olive-oil and diffused through one ounce of prepared lard.

QUESTIONS AND EXERCISES.

692. How is Aconitia prepared ?
693. Give the strengths of the official preparations of Atropia.
694. Describe the properties of atropia.
695. What is the active principle of stramonium ?
696. Mention pharmacopœial substances containing *berberia* and *berberia* respectively.
697. Give the characters of *berberia*.
698. In what does *nectandria* differ from *berberia* ?
699. Mention the characteristics of *conia*.
700. What is the active principle of *Ipecacuanha* ?
701. Name the alkaloid of Tobacco.
702. Give the properties of the alkaloid of Calabar Bean.
703. What are the sources of *piperia* ?
704. Whence is theia obtained ?
705. Describe the preparation of *Veratria*.
706. State the properties of *veratria*.

BITTER OR TONIC SUBSTANCES, ETC.

The following articles employed medicinally in such forms as Decoction, Extract, Infusion, Tincture, etc., contain active principles which have not yet been thoroughly examined. Some of these principles have been isolated, and a few have been obtained in the crystalline condition ; but their constitution has not been sufficiently well made out to admit of the classification of the bodies either among alkaloids, glucosides, acids, or other well-marked principles.

Andrographis Caules et Radix,
P. I., from *Andrographis*
paniculata, Kariyat.
Anthemidis flores.
Asclepias tuberosa, *Pleurisy-*
root (Asclepedin).
Aurantii cortex (Hesperidin).

Azadirachtæ Cortex et Folia,
P. I., from *Azadirachta in-*
dica, Nim or Margosa.
Baptisia tinctoria, wild indigo.
Alkaloid (?). B. L. Smedley,
Greene.

- Bonducella semina*, P. I., from *Cesalpinia* (*Guilandina*) *bonducella*, Bonduc seeds or Nicker nuts.
- Buchu folia*.
- Calotropis* Cortez, P. I., from *Calotropis procera* and *C. gigantea*, Mudar.
- Canella alba cortex*.
- Cascarilla cortex* (*Cascarillin*, $C_{12}H_{11}O_4$).
- Caulophyllum thalictroides*, Blue cohosh. Alkaloid (?).
- Cimicifuga* (*Actea*) *racemosa* (*Cimicifugin* ?).
- Cusparia cortex*, Angostura bark (*Cusparin*).
- Cypripedium pubescens* (*Cypripedin*).
- Euonymus Europæus* (*Euonymin*), Wahoo bark.
- Gulancha* (*Tinospora radix et caules*, P. I.).
- Gynocardia semina*, from *Gynocardia odorata* (*Chaulmúgra*), P. I.
- Hydrocotyles folia*, P. I., from *Hydrocotyle asiatica*, Indian pennywort.
- Iris versicolor*, *Blue Flag* (*Iridin* or *Irisin*).
- Lactuca*. The milk juice dried yields *lactucarium* (*Lactucin*, etc.).
- Lupulus*.
- Matica folia*.
- Quassia lignum* (*Quassin*, $C_{10}H_{12}O_3$).
- Serpentaria radix*.
- Soyimida Cortex*, P. I., Rohunbark, from *Soyimida febrifuga*.
- Taraxaci radix* (*Taraxacin*).
- Toddalia radix*, P. I.
- Veronica virginica* (*Leptandrin*).

AMYLACEOUS AND SACCHARINE SUBSTANCES.

STARCH.

Formula $C_6H_{10}O_5$; possibly $C_{72}H_{120}O_{60}$.

Processes.—Rasp or grate, or, with a knife, scrape a portion of a clean raw potato, letting the pulp fall on to a piece of muslin placed over a small dish or test-glass, and then pour a slow stream of water over the pulp; minute particles or granules of starch pass through the muslin and sink to the bottom of the vessel, fibrous matter remaining on the sieve. This is potato-starch. Even diseased potatoes furnish good starch by this method. Wheat-starch (*Amylum*, B. P.) may be obtained by tying up some flour in a piece of calico and kneading the bag in a slow stream of water flowing from a tap, the washings running into a deep vessel, at the bottom of which the white starch collects: the sticky matter remaining

in the bag is *gluten*. The *blue starch* of the shops is artificially coloured with smalt or indigo, to neutralize the yellow tint of recently washed linen; it should not be used for medicinal purposes. Starch dried in mass splits up into curious columnar masses, resembling the basaltic pillars of Fingal's Cave in Staffa, or those of the Giant's Causeway in the North of Ireland. The cause of the phenomenon, which may also be seen in grain-tin, is not conclusively known.

Gluten is the body which gives tenacity to dough and bread. It seems to be a mixture of vegetable fibrin, vegetable casein, and an albuminous matter termed gluten. These substances and gluten itself are closely allied; each contains about 16 per cent. of nitrogen. *Wheaten Flour* (*Farina Triticæ*, B. P.), contains about 72 per cent. of starch and 11 of gluten, as well as sugar, gum, fine bran, water, and ash. The compactness of barley, well seen in Husked or Pearl Barley (*Hordeum Decorticatum*, B. P.), is said to be due to the large amount of vegetable fibrin present. During germination the fibrin is destroyed; hence, probably, the cretaceous character of malt. *Oatmeal* (*Avenæ Farina*, U. S. P.) is very rich in albumenoid or flesh-forming constituents, containing nearly 16 per cent. *Sago* is granulated starch from the Sago Palm. *Tapioca* is granulated starch from the Bitter Cassava. The white translucent grains known as Rice are the husked seeds of *Oryza sativa*. Rice (*Oryza*) and the Flour of Rice, or Ground Rice (*Oryzæ Farina*), are official in the Pharmacopœia of India. Rice is quite a staple article of food in tropical countries. Ground rice resembles flour of wheat in composition, but contains from 85 to 90 per cent. of starch.

Mucilage of Starch.—Mix two or three grains of starch with first a little and then more water, and heat to the boiling-point; mucilage of starch (*Mucilago Amyli*, B. P.) results.

This mucilage or paste is not a true solution; by long boiling, however, a portion of the starch becomes dissolved. In the latter case the starch probably becomes somewhat altered.

Chemical Test.—To some of the mucilage add a little free iodine; a deep blue colour is produced.

This reaction is a very delicate test of the presence of either iodine or starch. The starch must be in the state of mucil-

age; hence in testing for starch the substance supposed to contain it must be first boiled with water. The solutions used in the reaction should also be cold, or nearly so, as the blue colour disappears on heating, though it is partially restored on cooling. The iodine reagent may be iodine-water or tincture of iodine. In testing for iodine its occurrence in the free state must be insured by the addition of a drop, or even less, of chlorine-water. Excess of chlorine must be avoided, or chloride of iodine will be formed, which does not colour starch.

The so-called *iodide of starch* scarcely merits the name of a chemical compound, the state of union of its constituents being feeble. Substances that attack free iodine remove that element from iodide of starch. The alkalies, hydrosulphuric acid, sulphurous acid, and other reducing agents destroy the blue colour.

Composition of Starch Granules.—Starch Granules consist mainly of *granulose*, soluble in cold water and giving an indigo colour with iodine, and *starch cellulose*, insoluble in water and giving with iodine a dirty yellow colour. The starch cellulose forms an external coating upon the granule and also exists mixed with the granulose inside the granule. If this coating be broken by mechanical means the continued application of cold water will remove all the granulose leaving the cellulose insoluble. By the action of diastase, ptyalin, and other ferments, and by other means, the granulose may be converted into sugar and dextrin, leaving the starch cellulose unacted upon.

Microscopical Examination of Starches.

All kinds of starch afford the blue colour with iodine, showing their chemical similarity. Physically, however, the granules of different starches differ from each other; hence a careful microscopical examination of any starch, or of any powder or vegetable tissue containing starch, enables the observer to state, with a high degree of probability, the source of the starch, either at once if he has much experience, or after comparing the granules in question with authentic specimens. A glance at the accompanying eight engravings* (figs. 41 to 48) of common starches will show to what extent

* By permission of Messrs. Longmans & Co., these engravings have been copied, with very few modifications, from the plates in two of the three volumes of the original edition of Pereira's "Materia Medica."

different starch-granules naturally differ in size, shape, general appearance, distinctness and character of the rugæ, and position of the more or less central point or hilum. While from different starches individual granules may be picked out which much resemble each other, the appearance of each starch as a whole is fairly characteristic; that is to say, each *group* of granules differs in one, two, or several characters from similar *groups* of granules of other starches.

A quarter-inch object-glass will commonly suffice for the microscopical observation of starch. A very little of the starch is mixed on a glass slide with a drop of water, a piece of thin covering-glass placed on the drop and gently pressed, so as to provide a very thin layer for observation. Instead of water, diluted spirit of wine, diluted glycerine, turpentine or other essential oil, Canada balsam, and other fluids may be used in cases where the markings or other appearances are not well defined. The illumination also of the granules may be varied, the light being reflected or transmitted, concentrated or diffused, white or coloured, polarized or plain. Polarized light is especially valuable in developing differences, and in intensifying the effects of obscure markings. By polarized light the granules of potato-starch appear as if traversed by a black cross; wheat starch granules and many others also peculiarly and characteristically influence polarized light. Distinctive characters will sometimes present themselves only when the granules are made to roll over in the fluid in which they have been temporarily mounted, or when the slide is gently warmed. Starches which have already been subjected to the influence of heat, partly, as in sago or tapioca, or almost entirely, as in bread, will of course differ in appearance from granules of the same starch before being dried, cooked, or torrefied. The characters of a starch will also somewhat vary according to the age and condition of the plant yielding it.

(For plates and descriptions of the characters of other starches occurring in plants used for medicinal purposes, the reader is referred to works on *Materia Medica*, and to the indexes of *Journals of Pharmacy*, as well as to general works and magazines on microscopy. For engravings of starch-granules *in situ*, *vide* Berg's "*Anatomischer Atlas*," published by Gaertner, Berlin.)

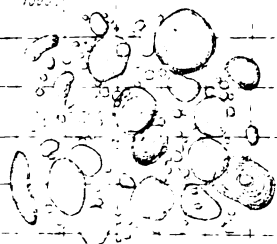
The student may place fair confidence in the accompanying lithographs, and in most of the published engravings of

STARCHES

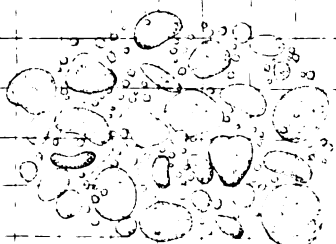
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W. J. F. 250 211

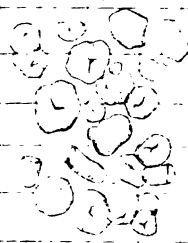
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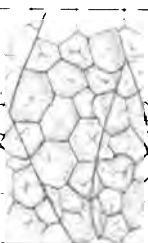
WHEAT (TRITICUM)



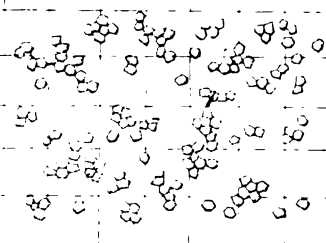
BARLEY (HORDEUM)



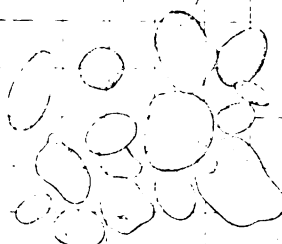
MAIZE (ZEA)



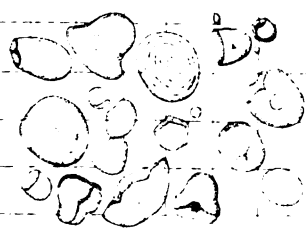
RICE (ORYZA)



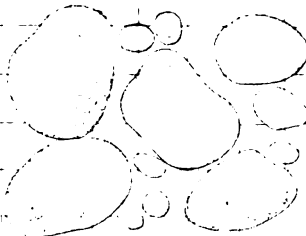
PEAR (PYRUS)



SWEET POTATO (IPOMOEA BATATAS)



TARO (COLOCASIA ESCULENTA)



POTATO (SOLANUM TUBEROSUM)

starch-granules; but in microscopical analyses of importance the worker should, if possible, himself obtain actual specimens of starches for comparison from the respective seeds, fruits, and other tissues.

Inulin ($C_6H_{10}O_5$) is a white powder apparently occupying the place of starch in the roots of many plants, especially those of the natural order *Compositae*. Twenty to forty-five per cent. has been obtained from elecampane (*Inula helenium*). It is also contained in the dahlia, colchicum, arnica, dandelion, chicory, etc. It is soluble in boiling water, nearly all being redeposited on cooling. Iodine turns it yellow. Long ebullition converts it into a kind of gum.

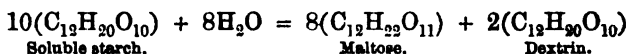
Lichenin ($C_6H_{10}O_5$) is a white starch-like powder largely contained in many lichens—Iceland "Moss," *Cetraria islandica*, and many others. It is soluble in boiling water, and the fluid gelatinizes on cooling. It may be precipitated from its aqueous solution by alcohol. With iodine it gives a reddish blue colour.

DEXTRIN.—Mix a grain or two of starch with about half a test-tubeful of cold water and a drop or two of sulphuric acid, and boil the mixture for a few minutes; no mucilage is formed, and the liquid, if sufficiently boiled, yields no blue colour with iodine; the starch has become converted into *dextrin* and sugar. Dextrin is also produced if starch is maintained at a temperature of about 320° F. for a short time. Dextrin is now largely manufactured in this way, and a paste of it used by calico-printers as a vehicle for colours; it is termed *British gum*. The change may also be effected by *diastase*, a peculiar ferment existing in malt. Mix two equal quantities of starch with equal amounts of water, adding to one a little ground malt, then heat both slowly to the boiling-point: the mixture without malt thickens to a paste or pudding; that with malt remains thin, its starch having become converted into dextrin and a sugar termed maltose.

Diastase is probably a mixture of the coagulable albumenoids. It is so named from *διάσρασις* (*diastasis*), *separation*, in allusion to the separation, or rather alteration, it effects among the constituent atoms of the molecule of starch. This function is shared by the saliva, pancreatic juice, bile, and

the intestinal and other juices. The function is destroyed when the albumenoids are coagulated by a temperature of from 176° to 178° F.

The action of diastase upon starch.—Diastase has scarcely any action upon unbroken starch granules. The granules must be ruptured by gelatinization with heat and moisture, or in some other way. When a solution containing diastase, such as a cold water infusion of malt, be allowed to act upon gelatinous starch or starch paste at 140° to 160° F., liquefaction occurs. It is possible to operate so that when liquefaction has taken place the solution shall give no reaction for sugar or dextrin. If this solution be concentrated and allowed to cool, a glistening white precipitate of soluble starch falls. Soluble starch is probably the result of the partial decomposition of the more complex molecule of granulose or gelatinous starch. The next step in the action of diastase upon gelatinous starch is the breaking down of the soluble starch molecule into dextrin and a sugar called maltose. At least ten dextrans are successively produced, each simpler than the one preceding it, the proportion of maltose being correspondingly increased. The dextrans first produced give a red or brown colour with iodine, while those last produced, and having a simpler molecule, give no colour with iodine. The final reaction may be expressed thus:—



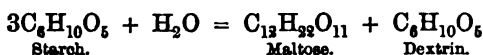
The dextrans are distinguished by their rotary power, their reducing action on cupric salts, and in other ways.

The action of dilute acids upon starch.—Dilute acids act upon gelatinous starch in the same way as diastase, except that the final product is glucose.

Malt (the word malt is said to be derived from the Welsh *mall*, soft or "rotten") is simply barley which has been softened by steeping in water, allowed to germinate slightly, and further change then arrested by the application of heat in a kiln. During germination the gluten breaks up and yields a glutinous substance termed vegetable gelatine, diastase, and other matters. To the vegetable gelatine is due much of the "body" of well malted and slightly hopped beer; it is precipitated by tannic acid; hence the thinness of ale (pale or bitter) brewed with a large proportion of hop or other materials containing tannic acid. A portion of the diastase

reacting on the starch of the barley converts it into dextrin, and, indeed, carries conversion to the further stage of grape-sugar, as will be explained immediately. The temperature to which the malt is heated is made to vary, so that the sugar of the malt may or may not be partially altered to a dark-brown colouring material: if the temperature is high, the malt is said to be *high-dried*, and is used in porter-brewing; if low, the product is of lighter colour, and is used for ale. The diastase remaining in malt is still capable of converting a large quantity of starch into dextrin and sugar (maltose); hence the makers or distillers of the various spirits operate on a mixture of malted and unmalted grain in preparing liquors for fermentation.

Extract of Malt is an evaporated infusion of malt. Taken with food its diastase aids in the conversion of starch into a variety of sugar termed maltose, and dextrin, and, *pro tanto*, assists enfeebled digestive powers.



As diastase loses this power at temperature above 150° F., that degree should not be exceeded in evaporating the infusion; indeed, if the dissolved albumenoid matters are to be retained, the evaporation should be conducted at 120° F. (For a method of estimating the strength of Extract of Malt, see a paper by Dunstan and Dimmock, *Pharm. Journ.*, March 8, 1879.)

Gum is a frequent constituent of vegetable juices, existing in large quantity in several species of Acacia. According to Fremy gum is a calcium salt, sometimes partially a potassium salt, of the *gummic* or *arabic* radical. The formula of gummic acid is said to be $\text{H}_2\text{C}_{12}\text{H}_{18}\text{O}_{10}$, H_2O . Gum differs from dextrin in yielding oxalic acid, but no mucic acid when oxidized by nitric acid. *Cerasin* or cherry-tree gum is a metagummate of calcium, an insoluble modification of acacia gum. *Bassorin*, *traganthin*, or *adraganthin* ($\text{C}_{12}\text{H}_{20}\text{O}_{10}$) is a form of gum which is insoluble in water, but absorbs large quantities of that liquid and forms a gelatinoid mass: it occurs largely in Tragacanth, combined, like arabin, with calcium. *Pectin*, or Vegetable Jelly ($\text{C}_{23}\text{H}_{40}\text{O}_{28}$, $4\text{H}_2\text{O}$), is the body which gives to expressed vegetable juices the property of gelatinizing: it forms the chief portion of Irish or Carrageen "Moss" (*Ohondrus crispus*). Ceylon "Moss"

(*Gracillaria lichenoides* and *G. confervoides*, P. I.) contains from one-third to three-fourths of vegetable jelly.

The mucilage of marsh-mallow root (*Althea officinalis*) and of linseed or common flax-seed (*Linum usitatissimum*) is a gum-like substance containing much mineral matter. It is the basis of the infusions termed *mallow-tea* and *linseed-tea*. Somewhat similar mucilage occurs in infusion of Bael: it is also largely yielded by the seeds of the Quince (*Cydonia vulgaris*), and by ordinary Elm-bark (*Ulm Cortex*, B. P.), as well as by the bark of the Red or Slippery Elm (*Cortex Ulmi Fulvæ*). *Salep*, the powdered dried tubers of many species of *Orchis*, contains a large quantity of such matter. *Squill* also. The Indian *Okra* (*Hibisci Capsule*, P. I., from *Hibiscus esculentus*) and *Ispaghul* or *Spogel* seeds (*Ispaghulæ semina*, P. I., from *Plantago ispaghula*) also appear to contain a considerable quantity.

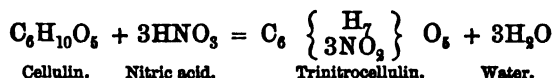
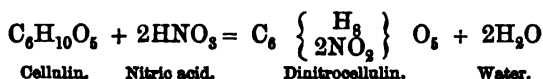
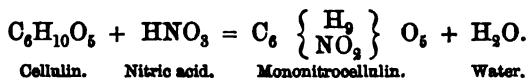
Isomerism. Allotropy. Polymorphism.

The composition of dextrin is represented by the same formula as that of starch, namely $C_6H_{10}O_5$; for it has the same percentage composition as starch. Inulin (p. 477) and cellulose (p. 482) have also a similar formula. There are many other bodies similar in centesimal composition, but dissimilar in properties; such substances are termed *isomeric* (from *ἴσος*, *isos*, equal, and *μέρος*, *meros*, part); and their condition is spoken of as one of *isomerism*. There is sometimes good reason for doubling or otherwise multiplying the formula of one of two *isomers*, *isomerides*, or isomeric bodies. Thus a molecule of olefiant gas (ethylene), the chief illuminating constituent of coal-gas, is represented by the formula C_2H_4 , while a molecule of amylene, an anæsthetic liquid hydrocarbon, obtained from amylic alcohol, though having the same percentage composition as olefiant gas, is represented by the formula C_5H_{10} ; for amylene, when gaseous, is about twice and a half as heavy as ethylene, and must contain, therefore, in each molecule, twice and a half as many atoms, for (Avogadro) these equal volumes must contain equal numbers of molecules; its formula is, consequently, constructed to represent those proportions. (Read again pages 29 to 59.) This variety of isomerism is termed *polymerism* (from *πολύς*, *polús*, many or much, and *μέρος*, *meros*, part). *Meta-stannic acid* (*vide* p. 278) is a polymeric variety, or *polymeride*, of stannic acid. An illustration of a second variety of iso-

merism is seen in the case of cyanate of ammonium and urea, bodies already alluded to in connection with cyanic acid. These and several other pairs of chemical substances have dissimilar properties, yet are similar not only in elementary composition and in the centesimal proportion of the elements, but also in the fact that each molecule possesses the same number of atoms. But the reactions of these bodies indicate the probable nature of their construction; and this is shown in their formulæ by the disposition of the symbols. Thus cyanate of ammonium is represented by the formula NH_4CNO , urea by $\text{CH}_4\text{N}_2\text{O}$. Such bodies are termed *metameric* (from $\mu\epsilon\tau\acute{\alpha}$, *meta*, a preposition denoting change, and $\mu\acute{\epsilon}\rho\omicron\varsigma$), and their condition spoken of as one of *metamerism*. Acetate of ethyl (p. 349) is metameric with butyric acid (p. 424), for they have the same percentage composition and their vapours have the same specific gravity, and each therefore might be represented by the formula $\text{C}_4\text{H}_8\text{O}_2$; but their properties warrant us in assuming that their atoms occupy different positions in the two molecules—justify us in giving $\text{C}_2\text{H}_5\text{C}_2\text{H}_3\text{O}_2$ as a picture of a molecule of acetate of ethyl, and $\text{HC}_4\text{H}_7\text{O}_2$ as a picture of a molecule of butyric acid. Acetate of methyl ($\text{CH}_3\text{C}_2\text{H}_3\text{O}_2$), propionic acid ($\text{HC}_3\text{H}_5\text{O}_2$), and formate of ethyl ($\text{C}_2\text{H}_5\text{CHO}_2$) are isomers of the metameric variety, or *metamerides*; also quinine and quinidine, cinchonine and cinchonidine, and many of the volatile oils, etc. The isomerism of starch and dextrin may be of a polymeric or of a metameric character; but we do not yet know which, and must therefore at present give them identical formulæ. Substances similar in composition and constitution, yet differing in properties, are termed *allotropic* ($\acute{\alpha}\lambda\lambda\omicron\varsigma$, *allos*, another, $\tau\rho\acute{o}\varsigma$, *tropos*, condition). Thus ordinary phosphorus, kept at a temperature of about 450°F ., in an atmosphere from which air is excluded, becomes red, opaque, insoluble in liquids in which ordinary phosphorus is soluble, oxidizes extremely slowly, and only ignites when heated to near 500°F . (red or amorphous phosphorus). A black allotropic variety of phosphorus is known. There are also three *allotropes* of carbon which are respectively crystalline, graphitic, and amorphous. Sulphur may be obtained in the viscous as well as in the hard, brittle condition. Another illustration of *allotropy* is seen in the varieties of tartaric acid, which have different optical properties, but otherwise are identical; they are in neither of the above-

mentioned states of isomerism, but are allotropic modifications of the same substance. Occasionally one and the same substance crystallizes in two distinct forms; its state is then described as one of *polymorphism* (πολὺς, *polús*, many, μορφή, *morphê*, form). Sulphur is polymorphous. It crystallizes by slow cooling in (1) prismatic crystals of sp. gr. 1·98, while in nature it occurs in (2) octahedra of sp. gr. 2·07. Melted and poured into water, sulphur takes up the form of (3) caoutchouc of sp. gr. 1·96. These differences warrant the statement that sulphur occurs in three distinct allotropic conditions. Possibly such conditions result from the association of different numbers of atoms in the molecule of the element, that is, allotropic bodies may simply be *physically polymeric*, or in some other way be mere *physical isomerides*.

Cellulin or *cellulose*, the woody fibre of plants, familiar, in the nearly pure state, under the forms of "cotton wool" (*Gossypium*, B. P., "hairs of the seed of various species of *Gossypium*"), paper, linen, and pith, is another substance isomeric, probably polymeric, with starch. Lignin is a closely allied body, lining the interior of woody cells and vessels. By the action of nitric acid of various strengths on cellulín, peroxide of nitrogen (NO_2) is substituted for one, two, or three atoms of hydrogen—mono-, di-, or trinitrocellulin being formed :—



Trinitrocellulin is highly explosive gun-cotton; dinitrocellulin is not sufficiently explosive for use instead of gunpowder; mononitrocellulin is scarcely at all explosive. The three movable atoms of hydrogen in cellulín may be displaced by bodies other than peroxide of nitrogen.

Dinitrocellulin (*Pyroxylin*, B. P.) may be prepared by the following official process :—Mix 5 fluid ounces of sulphuric

acid and 5 of nitric in an earthenware mortar, immerse 1 ounce of cotton-wool in the mixture, and stir it for three minutes with a glass rod, so that it is thoroughly and uniformly wetted by the acids. Transfer the cotton to a vessel containing a considerable volume of water, stir it rapidly and well with a glass rod, decant the liquid, pour more water upon the mass, agitate again, and repeat the affusion, agitation, and decantation until the washings cease to give a precipitate with chloride of barium. Drain the product on filtering paper, and dry in a water-bath.

Pyroxylin may also be made by soaking 7 parts of white filtering-paper, which has been washed in hydrochloric acid and dried, in a mixture of 140 parts of sulphuric acid (sp. gr. 1.82) and 70 of nitric acid (1.37) for 3 hours, and well washing the product (Guichard).

Mononitrocellulin and *trinitrocellulin* are insoluble in a mixture of alcohol and ether; *dinitrocellulin* or *pyroxylin* is soluble, the solution forming ordinary collodion (*Collodium*, B. P.). The official proportions are 1 oz. of pyroxylin dissolved in a mixture of 36 fl. oz. of ether and 12 of rectified spirit. After digesting for a few days, the liquid is decanted from any insoluble matter and preserved in a well-corked bottle. It is "a colourless highly inflammable liquid with ethereal odour, which dries rapidly upon exposure to the air, and leaves a thin transparent film, insoluble in water or rectified spirit." Flexible collodion (*Collodium Flexile*, B. P.) is a mixture of collodion (6 fl. oz.), Canada Balsam (120 grains), and castor-oil (1 fluid drachm).

QUESTIONS AND EXERCISES.

707. How is wheat-starch or potato-starch isolated?
708. Define gluten and glutin.
709. Enumerate the proximate principles of wheaten flour.
710. Is starch soluble in water?
711. Which is the best chemical test for starch?
712. Distinguish physically between the varieties of starch.
713. Into what compound is starch converted by heat?
714. What occurs when a mixture of starch and water is allowed to flow into hot diluted sulphuric acid?

715. If two equal amounts of starch with water be heated, one containing a small quantity of ground malt, what effects ensue?

716. Write a short article on the chemistry of "malting."

717. What is the nature of gum-arabic? and how is it distinguished from "British Gum"?

718. Explain *isomerism*, giving several illustrations.

719. Give examples of polymeric bodies.

720. State the formula of a body metameric with urea.

721. Define *allotropy* and *polymorphism*, giving illustrations.

722. What form of cellulin is official?

723. Mention the properties of the products of the action of nitric acid of various strengths on cellulin.

724. How is pyroxylin prepared?

SUGARS.

Sucrose or cane-sugar, $C_{12}H_{22}O_{11}$. Maltose, $C_{12}H_{22}O_{11}$.
 Lævulose or inverted sugar, $C_6H_{12}O_6$. Dextrose, glucose, or grape-sugar, $C_6H_{12}O_6$, H_2O . Lactose or milk-sugar, $C_{12}H_{22}O_{11}$, H_2O .

Artificial formation of Grape-sugar from Cane-sugar.—*Tests for Sugar.*—Dissolve a grain or two of common cane-sugar in water. To a portion of this solution placed in a test-tube add more water, two or three drops of solution of sulphate of copper, a considerable quantity of solution of potash or soda (enough to turn the colour of the liquid from a light to a dark blue), and heat the mixture to the boiling-point; no obvious immediate change occurs. To another portion of the syrup add a drop of sulphuric acid, and boil for ten or twenty minutes, then add the copper solution and alkali, and heat as before; a yellowish red precipitate of cuprous oxide (Cu_2O) falls. This test is exceedingly delicate.

The above reaction is due to the conversion of the *cane-sugar* ($C_{12}H_{22}O_{11}$) into *inverted sugar* or *lævulose*, $C_6H_{12}O_6$ (so called because its solution causes left-handed rotation of a ray of polarized light, cane-sugar having an opposite effect), and *grape-sugar*, $C_6H_{12}O_6$, H_2O , by the influence of the sulphuric acid, and to the reducing action of the inverted sugar and grape-sugar on the cupric solution. The formation of a

precipitate immediately, without the action of acid, shows the presence of the latter sugars—its formation only after ebullition with acid indicating, in the absence of starch or dextrin, cane-sugar. In this reduction process the sugar is oxidized and broken up into several substances; but the exact nature of the reaction has not been ascertained.

Dextrin also reduces the copper salt to suboxide, unless its solution is cold and very dilute. It does not, however, so act on a solution of cupric acetate acidified with acetic acid, while glucose produces with this liquid the usual red cuprous precipitate (Barfoed).

Cane-sugar or *sucrose* (*Saccharum Purificatum*, B. P.) is a frequent constituent of vegetable juices. Thus it forms the chief portion of cassia-pulp (*Cassia Pulpa*, B. P.), is contained in the carrot and turnip, but is most plentiful in the sugar-cane; much, however, is now obtained from the sugar-maple and beetroot. On evaporation of the juice, common *brown* or *moist sugar* crystallizes out; this by resolution, filtration through animal charcoal, evaporation to a strong syrup, and crystallization in moulds, yields the compact crystalline conical loaves known in trade as *lump-sugar*. From a slightly less strong syrup, slowly cooled, the crystals termed *sugar-candy* are deposited, white or coloured according to the colour of the syrup.

Maltose, $C_{12}H_{22}O_{11}$.—This crystallizable sugar is formed together with dextrin when diastase or dilute acids act upon starch. In the case of diastase it is the ultimate product, but the dilute acids may convert it into dextrose. It reduces cupric salts, but to a less extent than dextrose. It differs also from dextrose in its optical activity.

Inverted Sugar or *Lævulose* is uncrystallizable. It is found in the grape, fig (*Ficus*, B. P.), cherry, and gooseberry; both grape-sugar and inverted sugar in the strawberry, peach, plum, etc. Fruit sugar reduces cupric salts.

Grape-sugar, *glucose* (from γλυκὺς, *glucūs*, sweet) or *dextrose* is often seen in the crystallized state in dried grapes or raisins and other fruits; it is also the variety of sugar met with in diabetic urine. Its crystalline character is quite distinct from that of cane-sugar, the latter forming large four- or six-sided rhomboidal prisms, while grape-sugar occurs in masses of small cubes or square plates. Grape-sugar is also less soluble in water, but more soluble in alcohol than cane-sugar. Grape-sugar reduces cupric salts.

According to Fresenius, the percentage proportion of saccharine matter in the dried fig is 60 to 70, grape 10 to 20, cherry 11, mulberry 9, currant 6, whortleberry 6, strawberry 6, raspberry 4.

Lævulose is lævogyrate, while sucrose and glucose possess right-handed rotation; the latter twist a ray of polarized light from left to right, to an extent dependent on the amount of sugar present—a fact easy of application in estimating the amount of sugar in syrups or in diabetic urine.

Both cane-sugar, maltose, and grape-sugar yield alcohol and carbonic acid gas by fermentation, the cane-sugar probably always passing into grape-sugar before the production of alcohol commences.



In *bread-making*, some of the starch is converted into dextrin, and this into sugar by the ferment. The above action then goes on, the liberation of gas producing the *rising* or swelling of the mixture of flour, water, and yeast (dough)—the temperature to which the mass is subjected in the oven causing escape of most of the alcohol, and further expansion of the bubbles of carbonic acid gas in every part of the now spongy loaf. The carbonic acid gas gradually evolved when flour is worked up for bread with a mixture of dry bicarbonate of sodium and tartaric acid (best preserved by previous admixture with dried flour and a little carbonate of magnesium)—*baking-powder*—exerts similar influence. The least objectionable method of introducing carbonic acid gas, however, is that of Dauglish, whose patent *Aërated Bread* is made from flour by mere admixture with carbonic-acid water under pressure. On removal from the cylinder, the resulting dough expands by the natural elasticity of the imprisoned carbonic acid, and the bake-oven completes the process. The crumb of bread is official (*Mica Panis*, B. P.).

Milk-sugar or *lactose* ($\text{C}_{12}\text{H}_{24}\text{O}_{12}$) (*Saccharum Lactis*, B. P.), the sweet principle of the milk of various animals, is not susceptible of alcoholic or vinous fermentation; but it resembles grape-sugar in reducing an alkaline solution of copper with precipitation of suboxide. It is readily obtained from milk by adding a few drops of acid, stirring, setting aside for the *curds* to separate, filtering, evaporating the *whey* to a

small bulk, filtering again if necessary, and allowing to cool and crystallize. It usually occurs in trade "in cylindrical masses, two inches in diameter, with a cord or stick in the axis, or in fragments of cakes—greyish-white, crystalline on the surface and in its texture, translucent, hard, scentless, faintly sweet, gritty when chewed." It is soluble in 6 parts of cold and 3 of boiling water: slightly soluble in alcohol, insoluble in ether. Powdered milk-sugar is used in pharmacy as a vehicle for potent solid medicines. Milk-sugar is convertible, by the action of dilute acids, into "galactose" and "lactoglucose"; these may be reunited to form milk-sugar.

Action of Alkali on Sugar.—To a little solution of grape-sugar add solution of potash or soda, or solution of carbonate of potassium, and warm the mixture; the liquid is darkened in colour from amber to brown, according to the amount of sugar present.

Tests.—The copper-reaction, the fermentation process, and the effect of alkalies form three good tests of the presence of grape-sugar, and, indirectly, of cane-sugar. A piece of merino or other woollen material, previously dipped in a solution of stannic chloride and dried, becomes of a brown or black colour when dipped in a solution of glucose and heated to about 300° F. by holding before a fire.

Sugar from starch.—Boil starch with a little water and a drop of sulphuric acid as for dextrin, but continue the ebullition for several minutes: on testing a portion of the cooled liquid with iodine, and another portion with the heated alkaline solution of a copper salt as described on page 484, it will be found that the starch has nearly all become converted into a sugar—dextrose. Maltose is also formed, at first, but by the continued action of the acid is changed to dextrose. When made on a large scale, a warm (131° F.) mixture of starch and water of the consistence of cream is slowly poured into a boiling solution of one part of sulphuric acid in one hundred of water, the whole boiled for some time, the acid neutralized by chalk, the mixture filtered, the liquid evaporated to a thick syrup and set aside; in a few days it crystallizes to a granular mass resembling honey. In this opera-

tion a small quantity of dextrin remains with the glucose; but if the process be conducted under pressure, conversion, according to Manbré, is complete. Sugar made from the starch of rice, maize, etc., is now largely used as a partial substitute for malt in brewing. It is known as *patent sugar*, *saccharine*, *dextrine*, *maltose*, etc.

The sugar in fresh fruits is mainly cane-sugar; but by the action of the acid, or possibly of a ferment in the juice, it is gradually converted into *inverted sugar*, a variety differing from cane-sugar in being uncrystallizable, and in having an inverted or opposite influence on polarized light, twisting the ray from right to left (*lævogyrate*, having *lævo-rotation*—hence sometimes termed *lævulose*). Ripe Hips (*Rosæ Caninæ Fructus*, B. P.) contain 30 per cent. of such sugar, besides gum and acid malates and citrates. Fruit-sugar, as gathered in the form of syrup by bees, is probably a mixture of these two varieties. It is gradually altered to a crystalline or granular mass of grape-sugar, as seen in dried fruits, such as Raisins (*Uvæ*, B. P.) and the Prune (*Prunum*, B. P.), and in solidified honey (*Mel*, B. P.). This, the common form of grape-sugar, is *dextrogyrate*, and hence is sometimes termed *dextrose*, to distinguish it from *lævulose*. Honey often contains pollen, hairs, spores, the dust and dirt from the flowers, and various flocculent matters which cause it to ferment and yield mannite, alcohol, and acetic acid; hence for use in medicine it is directed (*Mel Depuratum*, B. P.) to be clarified by melting and straining while hot through flannel previously moistened with warm water. A mixture of clarified honey 80 per cent., acetic acid 10 per cent., and water 10 per cent. is official under the name of *Oxymel* (from *ὄξος*, *oxus*, acid, and *μέλι*, *meli*, honey). A similar mixture of honey with acetic acid containing the soluble portions of squill-bulbs (*Scilla*, B. P.) is known as *Oxymel of Squill* (*Oxymel Scillæ*, B. P.). Honey or cane-sugar are the bases of the official *Confections*.

"*Honey-dew*" is a viscid saccharine matter occasionally met with on the leaves of the lime, maple, black alder, rose and other trees. Sometimes it is sufficiently abundant to dry and fall on the ground, forming a veritable "shower of manna." It is a mixture of cane-sugar, inverted sugar, and dextrin.

Barley-sugar is made by simply heating cane-sugar till it

fuses, a change from the crystalline to the uncrystallizable condition occurring. *Treacle* (*Theriaca*, B. P.), *Molasses* or *Melasses* (from *Mel*, honey), chiefly results from the application of too much heat in evaporating the syrups of the sugar-cane; it is a mixture of cane-sugar with uncrystallizable sugar and colouring-matter. *Liquorice-root* (*Glycyrrhizæ Radix*, B. P.) contains a considerable quantity of uncrystallizable sugar.

Caramel.—Carefully heat a grain or two of sugar in a test-tube until it blackens; the product is *caramel* or *burnt sugar* (the *Saccharum Ustum* of pharmacy). It is used as a colouring agent for gravies, confectionaries, spirits, and similar materials.

Mannite ($C_6H_{14}O_6$).—Boil manna with alcohol, filter, and set aside; mannite separates in colourless shining crystals or acicular masses, to the extent of from 60 to 80 per cent. of the manna.

Manna, B. P., is "a concrete saccharine exudation from the stem of *Fraxinus Ornus* and *F. rotundifolia*; it is obtained by making incisions in the stem of the trees." It occurs in "stalactiform pieces from one to six inches in length, and one or two inches in width, uneven, porous, and friable, curved on one side, of a yellowish-white colour, with a faintly nauseous odour, and a sweetish taste." Mannite is also met with in celery, onions, asparagus, certain fungi and sea-weeds, occurs in the exudations of apple- and pear-trees, and is produced during the viscous fermentation of sugar.

Mannite is an alcohol, the radical of which is sexivalent (C_6H_9)^{vi}6HO (Wanklyn). It is closely related to the sugars, glucose becoming mannite by action of nascent hydrogen:—



Indeed glucose itself is probably an alcohol of another radical (C_6H_9)^{vi}6HO. Mannite does not undergo vinous fermentation in contact with yeast. It is soluble in 5 times its weight of cold water.

Mucic acid ($H_3C_6H_8O_8$) and *Saccharic Acid* ($H_2C_6H_8O_8$) are two isomeric bodies formed by the action of dilute nitric acid on sugar and mannite.

QUESTIONS AND EXERCISES.

725. How are cane-sugar and grape-sugar analytically distinguished?

726. Mention the chief sources of cane-sugar.

727. Describe the methods of extracting and purifying cane-sugar.

728. Give chemical explanations of the different processes of bread-making.

729. How is milk-sugar obtained? and in what respects does it differ from other sugar?

730. By what process may starch be entirely converted into sugar?

731. What is the difference between fruit-sugar and honey?

732. What is Oxymel?

733. Describe the effect of heat on cane-sugar.

734. Describe the source and character of manna.

735. Give the latest view of the constitution of mannite.

736. Whence are mucic and saccharic acids obtained?

THE GLUCOSIDES.

Source.—The Glucosides are certain proximate vegetable principles which, by ebullition with dilute acid, or other method of decomposition, take up the elements of water and yield glucose, accompanied by a second substance, which differs in each case according to the body operated on. Several of the glucosides which are of pharmaceutical interest will now be considered. Tannin, or tannic acid, is also a glucoside; it has been described among the acids.

There are indications that glucosides may be regenerated from the bodies into which they are converted by heat.

Note on Nomenclature.—The first syllable of the names of glucosides and neutral principles generally are commonly given in allusion to origin; the last syllable is *in*, which sufficiently distinguishes them as a class.

AMYGDALIN ($C_{20}H_{27}NO_{11}, 3H_2O$).—This body, obtained by Robiquet and Boutron-Charlard in 1830, was the first discovered glucoside (Liebig and Wöhler, 1837). It is a white crystalline substance, existing in the bitter (*Amygdala Amara*, B. P.) but not in the sweet almond (*Amygdala Dulcis*, B. P.).

About 2 per cent. is readily extracted by strong alcohol from the cake left when the fixed oil has been expressed from bitter almonds. From the concentrated alcoholic solution ether precipitates the amygdalin.

Make an emulsion of two or three sweet almonds by bruising and rubbing them with water, and notice that it has no odour of essential oil of bitter almonds; add a grain or two of amygdalin, an odour of essential oil of bitter almonds is at once developed. Bruise two or three bitter almonds and rub with water; the volatile oil is again developed.

The source of the hydride of benzoyl, or essential oil of bitter almonds, in these reactions is the amygdalin, which, under the influence of *synaptase* or *emulsin* (a nitrogenous, casein-like, ferment existing in both bitter and sweet almonds), splits up into the essential oil, hydrocyanic acid, and glucose:—



As each molecule of amygdalin yields one of hydrocyanic acid, a simple calculation shows that 17 grains (mixed with emulsion of sweet almonds) will be required to form one grain of real hydrocyanic acid, a quantity equivalent to 50 minims of the dilute hydrocyanic acid of the British Pharmacopœia. The hydrocyanic acid is probably in chemical combination with the oil.

Test.—The reaction between *synaptase* and amygdalin is applicable as a test of the presence of one by the addition of the other, even when mixed with much organic matter.

Cherry-Laurel water (*Aqua Laurocerasi*, B. P., by distillation with water from *Laurocerasi Folia*, B. P.) contains hydrocyanic acid derived from a reaction similar to, indeed probably identical with, that just described, for bitter-almond oil is simultaneously produced. But the proportion of amygdalin or analogous body in cherry-laurel leaves is most variable; hence the strength of the water is highly uncertain.

Cortex Pruni Serotinæ, U. S. P.—The recently dried Wild-Black-Cherry Bark also furnishes by distillation an essential oil and hydrocyanic acid. Quince-seeds also (*Cydonia vulgaris*). The Wild Black Cherry, *Cerasus* (*Prunus*) *Virginiana*, contains amygdalin.

Caution.—Essential oil of almonds is of course highly poisonous. The purified oil or hydride of benzoyl is almost innocuous; it is obtained on distilling the crude oil with milk of lime and ferrous chloride and drying the product by shaking with fused chloride of calcium. The so-called "artificial oil of bitter almonds" or nitrobenzol [$C_6H_5(NO_2)H$], when taken in quantity, has been known to produce death. The presence of nitrobenzol in oil of bitter almonds is detected by adding a little of the oil to a mixture of zinc and diluted sulphuric acid, shaking well, setting aside for an hour or two, filtering off the clear liquid, and adding a little chlorate of potassium; a violet colour (actual mauve) is produced. Or the specimen may be shaken with bisulphite of sodium to fix the essential oil and then with ether, which dissolves out, and on evaporation will yield, the nitrobenzol.

ARBUTIN ($C_{24}H_{32}O_{14}$, H_2O) is contained in the leaves of *Arctostaphylos uva ursi* and *Chimaphila umbellata*. It is a bitter neutral body occurring in acicular crystals, and resolvable by acids into *hydrokinone* ($C_6H_6O_3$) and glucose, and by gentle oxidation into *kinone* ($C_6H_4O_3$) and formic acid. *Eriocolin* ($C_{34}H_{46}O_{21}$) is another bitter glucoside in bearberry-leaves.

CATHARTIC ACID.—"The glucoside acid that now is known to confer on the Senna of Alexandria (*Senna Alexandrina*, B. P.) and of Tinnevely (*Senna Indica*, B. P.) its purgative property, has been named by its discoverers (Dragendorff and Kubly) Cathartic acid. Its formula has been stated as $C_{180}H_{192}N_4SO_{82}$, which, if true, accounts for its extreme instability. It is insoluble in water, strong alcohol, and ether, but enters readily into watery solution when combined with alkaline and earthy bases, in which state it exists in senna. Its ammonium salts give brownish flocculent precipitates with salts of silver, tin, mercury, copper, and lead. Antimonial salts, tannin, yellow and red prussiates have no effect upon it. Alkalies, aided by heat, act destructively upon it. Boiled with a mineral acid it splits into a peculiar kind of glucose and an acid that has been named Cathartogenic; its formula is said to be $C_{133}H_{116}N_4SO_{44}$. The natural cathartate occurring in senna is prepared by partially precipitating by strong spirit a watery infusion of senna, concentrated to a syrupy state by evaporation *in vacuo*. The filtrate is now treated with a much larger bulk of absolute alcohol, and the precipitate thus obtained is purified by repeated solution in water and precipitation by alcohol. To obtain the

pure acid, advantage is taken of its colloidal properties; the crude cathartate is dissolved in moderately strong hydrochloric acid, and subjected to dialysis on a diaphragm of parchment paper. The minimum dose of this pure acid was found to be about $1\frac{1}{2}$ grain, which caused several stools with decided griping.

"The cathartic combinations that I have made are, the cathartate of ammonium, prepared from cathartate of lead by my original process, and the mixed cathartates, prepared according to Dragendorff's method as modified by myself. Of the former nearly pure salt, I have found $3\frac{3}{4}$ grains to purge fairly as to amount, but slowly as to time, and with considerable griping. Of the latter, $7\frac{1}{2}$ grains purged violently with much griping and sickness, which continued through the greater part of the day. It obviously would be improper to combine senna with any of its metallic precipitants, should such be desired, which is not likely. It is here satisfactory to observe that the cathartate of magnesium is soluble, and that the old-fashioned black draught agrees with new-fashioned science" (Groves).

Buckthorn-juice (*Rhamni Succus*, B. P.) owes its cathartic properties to a substance apparently identical with cathartic acid.

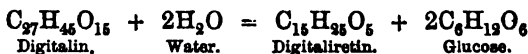
COLOCYNTHIN ($C_{56}H_{84}O_{23}$?).—This substance is the active bitter and purgative principle of colocynth-fruit (*Colocynthis Pulpa*, B. P.): it is soluble in water and alcohol, but not in ether. By ebullition with acids it furnishes glucose and a resinoid body.

CONVOLVULIN.—See JALAPIN.

COTOIN ($C_{23}H_{18}O_6$) appears to be the chief active principle of Coto bark, a Bolivian remedy for diarrhoea.

DAPHNIN ($C_{31}H_{34}O_{10}$) is the crystalline glucoside of the bark of *Daphne mezereum* (*Mezerei Cortex*, B. P.). Boiled with dilute acids it yields *daphnetin* ($C_{19}H_{14}O_9$) and glucose. The acrid principle of mezereum is resinoid.

DIGITALIN ($C_{27}H_{46}O_{15}$, Kosmann; $C_6H_8O_2$, Schmiedeberg).—This is an active principle of the Foxglove (*Digitalis*, B. P.). Boil a grain of digitalin (*Digitalinum*, B. P.) with dilute sulphuric acid for some time; flocks of *digitaliretin* ($C_{15}H_{25}O_5$) separate, and glucose may be detected in the liquid.



Properties.—Digitalin occurs "in porous mammillated masses or small scales, white, inodorous, and intensely bitter, readily soluble in spirit, but almost insoluble in water and in pure ether, dissolves in acids, but does not form with them neutral compounds; its solution in hydrochloric acid is of a faint yellow colour, but rapidly becomes green. It leaves no residue when burned with free access of air. It powerfully irritates the nostrils, and is an active poison." According to Pettenkofer "an intense red colour is produced if a trace of digitalin dissolved in water is mixed with a weak aqueous solution of inspissated bile and sufficient oil of vitriol added to raise the temperature to 158° F." Moistened with sulphuric acid and the liquid exposed to the vapour of bromine, a violet colour is produced.

Process.—The official process for the preparation of digitalin consists in dissolving the glucoside out of the digitalis-leaf (*Digitalis Folia*, B. P.) by alcohol, recovering the alcohol by distillation, dissolving the residue in water by the help of a small quantity of acetic acid, removing much of the colour from the solution by animal charcoal, neutralizing most of the acetic acid by ammonia, precipitating the digitalin by tannic acid (with which it forms an insoluble compound), washing the precipitate, rubbing and heating it with spirit and oxide of lead (which removes the acid in the form of insoluble tannate of lead), again decolourizing by animal charcoal, evaporating to dryness, washing out impurities still remaining by ether, and drying the residual digitalin. In this form digitalin is uncrystallizable.

Pure Digitalin (?).—On treating commercial digitalin with chloroform only an inert substance remains undissolved. The solution yields pure digitalin on evaporation; it may be crystallized from spirit in radiating needles (Nativelle.) The therapeutic effect of the pure substance is identical with the preparations of digitalis, but, as might be expected, more constant in its action, and, of course, intensely powerful.

Digitoxin ($\text{C}_{31}\text{H}_{53}\text{O}_7$) is a highly poisonous substance extracted from Foxglove by Schmiedeberg. The same chemist regards commercial digitalin from Foxglove-seeds as composed of three glucosides, namely, pure *digitalin* ($\text{C}_t\text{H}_8\text{O}_3$), *digitonin* ($\text{C}_{31}\text{H}_{52}\text{O}_{17}$) (closely allied to saponin), and *digitalein*.

ELATERIN ($C_{20}H_{28}O_6$).—Boil elaterium (*Elaterium*, B. P.), the dried sediment from the juice of the squirting-cucumber fruit (*Ecbalii Fructus*, B. P.), in a small quantity of spirit of wine, and filter; fibrous and amylaceous matter remain insoluble, while elaterin and resin are dissolved. The filtrate, concentrated and poured into a warm solution of potash, yields, on cooling, crystals of elaterin, resin being retained by the alkali. It is purified by recrystallization from spirit. Boil elaterium in dilute sulphuric acid for an hour or two, filter, and test the clear liquid for glucose; a reddish precipitate of cuprous oxide falls. This reaction is readily obtained with elaterium, but not always with elaterin; hence probably the latter is not a true glucoside. Walz states that elaterium also contains *prophetin*, a true glucoside.

Elaterin is the active principle of the so-called elaterium. Elaterium occurs "in light friable slightly incurved cakes, about one line ($\frac{1}{12}$ inch) thick, greenish-grey, acrid and bitter; fracture finely granular." Good specimens of this drug should yield, according to the British Pharmacopœia, not less than 20 per cent. of elaterin by the above process. Elaterium adulterated with chalk and other substances was formerly occasionally met with.

According to the experience of Flückiger and Hanbury the best method of obtaining elaterin is to exhaust elaterium with chloroform and then to add ether to the chloroform, when crystalline elaterin is precipitated. It should be washed with a little ether and crystallized from chloroform. When pure it occurs in hexagonal scales or prisms.

Test.—A little is placed in a watch-glass with a drop or two of liquefied carbolic acid, and then two or three drops of strong sulphuric acid: a carmine colour is developed. (Lindo).

GENTIOPICRIN, or **GENTIAN-BITTER** ($C_{20}H_{30}O_{12}$), the neutral crystalline principle of the root of *Gentiana lutea* (*Radix Gentianæ*, B. P.). It is soluble in water and weak spirit. Alkalies decompose it. Dilute acids convert it into *gentiogenin* and glucose. Gentian root also contains a crystalline acid ($HC_{14}H_9O_6$) termed *gentianic*, or *gentisic acid*, or *gentisin*.

Fused [potash, etc., gives] with the latter an acid ($C_7H_6O_4$) which has also, unfortunately, been called gentisic acid.

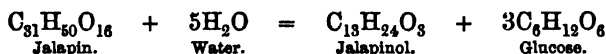
GLYCYRRHIZIN ($C_{24}H_{36}O_9$, Gorup-Besanez).—Liquorice-root (*Glycyrrhizæ Radix*, B. P.), in addition to uncrystallizable sugar, contains 3 or 4 per cent. of a sweet substance, *glycyrrhizin*, which, when boiled with hydrochloric acid or dilute sulphuric acid, yields a resinoid bitter body, *glycyrretin*, and an uncrystallizable sugar resembling glucose. Glycyrrhizin is only slightly soluble in cold water, hence is almost wholly excluded from the official evaporated infusion! (*Extractum Glycyrrhizæ*, B. P., and *Extractum Glycyrrhizæ Liquidum*, B. P.), but is present in considerable quantity in the evaporated decoction (*Stick Liquorice*, *Spanish Liquorice*, or *Solazzi Juice*). The tropical substitute for liquorice is the root of *Abrus precatorius* or *Indian Liquorice* (*Abri Radix*, P. I.), which also apparently contains glucose and glycyrrhizin. Glycyrrhizin has considerable power of disguising nauseous flavours. Roussin refers the sweet taste of liquorice not to pure glycyrrhizin but to a combination of glycyrrhizin with alkalies, and states that ammoniacal glycyrrhizin has exactly the sweetness of liquorice-root. The formula of this *glycyrrhizate of ammonium* is said by Habermann to be $(NH_4)_3C_{44}H_{60}NO_{18}$. Sestini finds that the glycyrrhizin of liquorice-root is chiefly glycyrrhizate of lime.

GUAIACIN.—Resin of guaiacum (*Guaiaci Resina*, B. P.), an exudation from the wood (*Guaiaci Lignum*, B. P.) of *Guaiacum officinale*, is probably a mixture of several substances, among which are *Guaiaretic* or *Guaiaretinic acid* ($C_{20}H_{28}O_4$) (Hlasiwetz), *Guaiaconic acid* ($C_{38}H_{40}O_{10}$) (Hadelich), and *Guaiacin*, a glucoside. On boiling guaiacum-resin with dilute sulphuric acid for some time, glucose is found in the liquid, a green resinous substance (*guaiaretin*) remaining insoluble (Kosmann). Most oxidizing agents, and even atmospheric air, especially under the influence of certain organic substances, produce a blue, then green, and finally a brown colour when brought into contact with an alcoholic solution of guaiacum-resin.

These effects are said to be due to three stages of oxidation (Jonas). They may be observed on adding the solution to the inner surface of a paring of a raw potato.

HELLEBORIN ($C_{36}H_{48}O_6$) and HELLEBOREIN ($C_{36}H_{44}O_{16}$) are crystalline glucosides occurring in the roots of Black Hellebore (*Helleborus niger*), or Christmas Rose, and Green Hellebore (*H. viridis*), ranunculaceous herbs.

JALAPIN ($C_{31}H_{50}O_{16}$) and CONVULVULIN ($C_{34}H_{56}O_{16}$).—According to Keyser and Meyer, jalap-resin contains two distinct substances—convulvin, chiefly obtained from Mexican male jalap (*Ipomæa orizabensis*), and jalapin, most largely contained in the true jalap (*Ipomæa purga*); the former is soluble in ether, the latter insoluble. Boil jalap-resin with dilute sulphuric acid for some time and filter; a substance, which is probably a mixture of *jalapinol* ($C_{13}H_{24}O_3$) and *convulvulinol* ($C_{16}H_{30}O_3$), separates; and glucose may be detected in the clear liquid. (It is to be regretted that the authors transpose the above names, terming the old well-known jalapin convulvin.)



Jalapic acid.—This is contained in the portion of jalap-resin soluble in ether. It may also be obtained from jalapin by ebullition with alkalis:—



Jalap-resin (*Jalapæ Resina*, B. P.) is obtained by digesting and percolating jalap-tubercles (*Jalapa*, B. P.) with spirit of wine, adding a little water, distilling off the spirit, pouring away the aqueous portion, which contains much saccharine matter, and washing and drying the residual resin. The tincture is sometimes decolourized by animal charcoal, and the evaporated product sold as “jalapin.”

Jalap-resin is insoluble in oil of turpentine; common resin, or rosin, soluble. If the presence of the latter is suspected, the specimen should be powdered, digested in turpentine, the

mixture filtered, and the filtrate evaporated: no residue, or not more than yielded by the turpentine itself, should be obtained.

Tampico Jalap, from *Ipomœa simulans*, yields a resin which apparently is chiefly convolvulin, but sometimes contains jalapin; for a sample obtained by Hanbury was entirely soluble in ether, and another extracted by Umney was almost wholly soluble, while Evans purified some, half only of which was soluble.

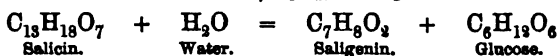
The *Kaladana resin*, or *Pharbitisin* of India (from *Pharbitis Nil*, P. I.), is a cathartic analogous to, if not identical with, resin of jalap.

PICROTOXIN ($C_8H_6O_3$, Oppermann; $C_{13}H_{14}O_5$, Pelletier and Couerbe) is a crystalline bitter poisonous principle (*πικρὸς*, *picros*, bitter, and *τοξικὸν*, *toxicon*, poison) occurring in *Cocculus indicus*, the dried fruits of *Anamirta cocculus*, P. I. Ludwig regarded it as a glucoside; but its constitution is not yet satisfactorily ascertained.

SALICIN ($C_{13}H_{18}O_7$).—This substance is contained in and easily extracted from willow-bark.

Tests.—1. To a small portion of salicin placed on a white plate or dish add a drop of strong sulphuric acid; a deep red colour is produced.

2. Boil salicin with dilute sulphuric acid for some time; it is converted into *saligenin* ($C_7H_8O_2$) and glucose.

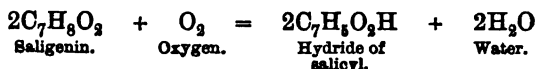


Examine a portion of the solution for grape-sugar by the copper test.

3. To another portion of the liquid, carefully neutralized, add a persalt of iron; a purplish-blue colour is sometimes produced, due to the reaction of the saligenin and the ferric salt. The saligenin is, however, so rapidly decomposed by acids into *saliretin* (C_7H_6O) and water, that this reaction is almost valueless as a test. Saligenin may readily be obtained by action of synaptase on salicin.

4. Heat a mixture of about 1 part of salicin, 1 of red chromate of potassium, $1\frac{1}{2}$ of sulphuric acid, and 20 of water in a

test-tube; a fragrant characteristic odour is evolved, due to the formation of hydride of salicyl ($C_7H_5O_2H$), an essential oil identical with that existing in meadow-sweet (*Spiræa ulmaria*) and in heliotrope.



SANTONIN ($C_{15}H_{18}O_3$).—This substance is, apparently, the anhydride of a weak acid (Hesse) insoluble in ammonia, but forming a soluble calcium salt. Indeed by boiling santonin for twelve hours with baryta water, Cannizarro has obtained a salt from which hydrochloric acid separates *santonie acid* ($C_{15}H_{20}O_4$). From a solution of santonate of calcium the santonin is precipitated by acids. Boiled for some time with dilute sulphuric acid it yields 87 per cent. of an insoluble resinous substance (*santoniretin*) and glucose (Kosmann). *Santonin* (*Santoninum*, B. P.) is official in the British Pharmacopœia: it is soluble in an aqueous solution of twice its weight of carbonate of sodium.

Process.—The process for its preparation consists in boiling *santonica* (the unexpanded flower-heads of an undetermined species of *Artemisia*) with milk of lime (whereby santonate of calcium is formed), straining, precipitating the santonin or santonie acid by hydrochloric acid, washing with ammonia to remove resin, dissolving in spirit and digesting with animal charcoal to get rid of colouring-matter, setting the spirituous solution aside to deposit crystals of santonin, and purifying by recrystallization from spirit (Mialhe).

Test.—To highly dilute solution of perchloride of iron add an equal bulk of concentrated sulphuric acid. To this reagent add the santonin, or powder or substance suspected to be santonin, and cautiously apply heat. A red, purple, and finally violet colour is produced (Lindo). *Santonin* added to warm alcoholic solution of potash yields a violet-red colour.

SAPONIN ($C_{12}H_{20}O_7$?) is a peculiar glucoside occurring in Soapwort, the root of the common Pink, and many other plants: its solution in water, even though very dilute, froths like a solution of soap. Pereira considered *emilacin* (*salseparin* or *parallin*), one of the principles of the supposed activity of *Sarsaparilla* (*Sarsæ Radix*, B. P.), to be closely allied to, if not identical with, saponin. According to Klunge ('Pharmacographia'), *parallin*, by action of acids, yields *parigenin*;

but his researches are not yet completed. The aqueous solutions of parallin froth when shaken.

Saponin is also met with in the root of *Polygala Senega* (*Senegæ Radix*, B. P.), though the active principle of senega is said to reside in *polygalic acid*, probably a glucosidic derivative of saponin.

SCAMMONIN ($C_{34}H_{56}O_{16}$).—Boil resin of scammony (*Scammonie Resina*, B. P.) with dilute sulphuric acid for some time; glucose may then be detected in the liquid, a resinous acid termed *scammoniol* ($C_{14}H_{18}O_3$?) being produced at the same time.

Natural scammony (*Scammonium*, B. P.) is an exudation from incisions in the living root (*Scammonie Radix*, B. P.) of *Convolvulus Scammonia*. It contains from 10 to 20 per cent. of gum, and, therefore, when rubbed up with water, gives an emulsion. "Ether removes from 80 to 90 per cent. of resin" (B. P.). The official resin of scammony contains no gum, and therefore gives no emulsion when rubbed up with water. It is made by digesting the root in spirit, adding water, distilling off the alcohol, and washing the residual resin with hot water till free from gum. There seems to be little or no chemical difference between the extracted resin and the resin of the exuded scammony.

Resin of scammony is soluble in all proportions in ether. Spirgatis states that it is identical with the resin of Mexican Male Jalap, which also is soluble in ether. Sulphuric acid slowly reddens it. It is said to be liable to adulteration with resin of true jalap, guaiacum-resin, and common rosin. Resin of true jalap is insoluble in ether, guaiacum-resin is distinguished by the colour-tests mentioned under GUALACIN, and rosin by the action of sulphuric acid.

SCILLITIN.—Schroff believes the bitter principle of the squill-bulb (*Scilla*, B. P.) to be a glucoside. Merok has extracted substances which he has termed *scillipicrin* and *scillitoxin*. But no definite crystalline principle has yet been obtained. Squill contains a large quantity of mucilage.

The bulbous root of *Crinum asiaticum* is official in the Pharmacopœia of India (*Crini Radix*, P. I.) as a substitute for squill. It has not been chemically investigated.

QUESTIONS AND EXERCISES.

737. Define glucosides, and mention those of pharmaceutical interest.

738. Draw out an equation illustrative of the development of Oil of Bitter Almonds.

739. How much pure amygdalin will yield one grain of real hydrocyanic acid?

740. To what does Cherry-Laurel water owe activity? Is the preparation trustworthy?

741. Mention the active principle of Senna.

742. By what process is the glucoside of the purple foxglove prepared?

743. State the circumstances under which Guaiacum-Resin and Jalap-resin yield glucose.

744. Mention a test for guaiacum-resin.

745. How may the adulteration of jalap-resin by rosin be detected?

746. Enumerate the tests for Salicin.

747. How is Santonin officially prepared?

748. Name sources of saponin.

749. What is the difference between Scammony and Resin of Scammony?

750. How would you detect resins of turpentine, guaiacum, or jalap in resin of scammony?

ALCOHOL AND ALLIED BODIES.

ALCOHOL, OR HYDRATE OF ETHYL.

Formation of Alcohol.—Ferment two or three grains of sugar by dissolving in a test-tubeful of water, adding a little yeast (*Cerevisia Fermentum*, B. P.), or a piece of the so-called German or dried yeast, and setting the whole aside for several hours in a warm place at a temperature of 70° or 75° F.; carbonic acid gas is evolved, and, if the tube be inverted in a small dish containing water, may be collected in the upper part of the tube and subsequently tested: the solution contains alcohol. If the experiment be made on larger quantities (four ounces of sugar, one of yeast, and a pint of water) the

fermented liquid should be distilled, one-half being collected, shaken with a little lime, soda, or potash to neutralize any acetic acid and decompose ethereal salts, and again distilled till one-half has passed over; the product is dilute spirit of wine. It may be still further concentrated or rectified by repeating this process of *fractional distillation*.

Fermentation.—The act of fermentation is commonly the result of or rather accompaniment of some vital action. *Alcoholic fermentation* would appear to be always attended by or to attend development of life and free multiplication of cellular structure. It follows the development of the fungus already referred to as constituting the chief active part of yeast, the *saccharomyces cerevisiæ*. In the presence of this fungus, with small quantities of phosphates and albumenoid matter, *glucose* is converted into alcohol and carbonic acid gas, together with small proportions of glycerin, succinic acid, and other substances. Yeast also contains a soluble ferment analogous to diastase, which is capable of converting sucrose into glucose. Therefore if yeast be used *sucrose* or cane-sugar may be converted into carbonic acid gas and alcohol, the soluble ferment first converting the sucrose into glucose.



Not more than 20 per cent. by weight of alcohol can be obtained in a fermenting fluid, for more than this proportion prevents fermentation.

Other kinds of fermentation, arising from the action of special ferments which have not received in all cases distinctive names, are the following. *Viscous* or *Mannitic fermentation*, which occurs when beer or saccharine juices, such as that of beet-root, become "ropy." Gum, mannite, and carbonic acid gas are produced. For *Lactic* and *Butyric fermentations* see Lactic acid. *Putrefactive fermentation* occurs when a liquid containing albumenoid matter is exposed to the air. Infusoria appear in the liquid, using up the dissolved oxygen, and the ferments of the genus *vibrio* are developed. These are protected from oxygen, which is fatal to them, by a thin surface layer crowded with bacteria—small rod-like organisms having powers of locomotion. The vibrionic action or putrefaction proceeds with evolution of sulphuretted

hydrogen, together with other gases having unpleasant odours and of complex chemical constitution. For *Acetic fermentation* see *Acetic acid*. For *Ammoniacal fermentation* see *Urine*.

Fermentation by certain soluble albumenoids. For the conversion of starch into sugar by diastase, see *Starch*; of amygdalin into benzoic aldehyd, hydrocyanic acid, and glucose by emulsin, see *Amygdalin*; of salicin into saligenin and glucose, see *Salicin*; of myronate of potassium into sulphocyanide of allyl, etc., by myrosin, see *Mustard*; of cane-sugar into grape-sugar by the soluble ferment in yeast, see the foregoing paragraphs.

Alcoholic fermentation.—The chief reaction results, as already stated, in the formation of alcohol and carbonic acid gas, though traces of several other substances are simultaneously produced (*vide* "Fousel Oil" in Index). By this reaction are formed the spirit of the various kinds of wine, beer, and liqueurs, such as Orange Wine (*Vinum Aurantii*, B. P.), made "by the fermentation of a saccharine solution, to which the fresh peel of the bitter orange has been added;" Sherry Wine (*Vinum Xericum*, B. P.), the fermented juice of the grape; Bay Rum, or Spirit of Myrcia (*Spiritus Myrciæ*, U. S. P.), prepared by distilling rum with leaves of *Myrcia acris*; and others.

Alcoholic drinks vary much in strength. Cider or apple wine, perry or pear wine, and good beer (ale and porter or stout) contain 4 to 6 per cent. of real alcohol; good light wines, both "red" and "white," 10 to 12 per cent.; good sherry and port, which are commonly "fortified," that is contain added spirit, 16 or 18 per cent.; while "spirits," (gin, rum, brandy, whisky, etc.) and "liqueurs," (ratafia, almond-flavoured; maraschino, cherry-flavoured; curaçoa, orange-flavoured; chartreuse, a composite flavoured liqueur, etc.) are "under-proof" or "over proof," terms explained in the next paragraph. The well known effects of these spirituous fluids on the animal system would appear to be due primarily to alcohol, and secondarily to ethereal derivatives of alcohols. Some owe a part of their effect to non-volatile substances, for beer from which all alcohol, etc., has been removed by ebullition still has a powerful influence on the human economy.

Varieties of Alcohol.—The weak spirit concentrated by distillation till it contains 84 per cent. by weight of pure alcohol is an ordinary article of trade; its specific gravity at 60° F.

is 0.8382. This is common *Spirit of Wine*, the *Spiritus Rectificatus* of the British Pharmacopœia. The official *Proof Spirit** (*Spiritus Tenuior*, B. P.) contains 49 per cent. by weight of alcohol, and is made by diluting 100 volumes of Rectified Spirit with water until the well-stirred product measures 156 volumes. Sixty volumes of water will be required for this purpose, the liquids occupying less bulk after than before admixture. In the language of the Excise authorities, the rectified spirit of the Pharmacopœia would be described as "56 per cent. over proof" (56 per cent. O. P.); that is, 100 volumes contain as much alcohol as is present in 156 volumes of proof spirit. Obviously, proof spirit may be made by diluting with water rectified spirit of any other strength than that mentioned above. Thus 100 fluid ounces of a spirit of "seventy over proof" may be diluted to 170, or the same quantity of a spirit of "fifty over proof" may be diluted to 150, and so on. The specific gravity of proof spirit at 60° F. is 0.920.

Empirical Formulas. Composition of Alcohol.—Alcohol, by quantitative analysis, is found to contain the elements carbon, hydrogen, and oxygen in the following proportions:—

Composition of Alcohol.

Carbon . . .	52.174 or ÷ 12 =	4.348 or 2.
Hydrogen . .	13.043 or ÷ 1 =	13.043 or 6.
Oxygen . . .	34.783 or ÷ 16 =	2.174 or 1.
100.000		

From the centesimal numbers a formula is obtained in the usual way. Thus, on dividing these figures by the atomic weights of the respective elements (C=12, H=1, O=16), and reducing the products to the simplest whole numbers, alcohol will be found to contain two atomic weights of carbon to every six of hydrogen and to every one of oxygen, and its possible or *empirical formula* to be C_2H_6O .

Constitution of Alcohol.—There is good reason to believe

* *Proof spirit* is so termed from the fact that in olden times a proof of its strength was supposed to be afforded by moistening a small quantity of gunpowder and setting light to the spirit; if it fired the powder it was said to be "over proof;" if not, "under proof." The weakest spirit that would stand this test was what we should now describe as of sp. gr. 0.920.

that alcohol is the hydrate of a basylous radical *ethyl* (C_2H_5 or Et); hence we derive the *rational formula* C_2H_5HO or $EtHO$.

Rational Formulæ.—Rational formulæ are deduced by (1) ascertaining how much of the substance will combine with, displace or play the part of, the atomic weight of a well-known element or radical. When this method cannot be applied, or in confirmation of it, processes of (2) reduction, (3) oxidation, (4) substitution, etc., are employed.

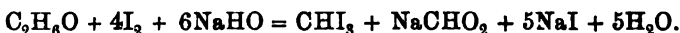
Salts of ethyl.—Alcohol is, then, a body analogous in constitution to hydrate of potassium (KHO); and there are other compounds of ethyl analogous in constitution to ordinary inorganic salts, such as those of potassium. The oxide of ethyl (Et_2O) is common ether; the nitrite of ethyl ($EtNO_2$) is the chief body which, dissolved in spirit of wine, constitutes "sweet spirit of nitre;" the acid sulphate of ethyl ($EtHSO_4$), or sulphethylic or sulphovinic acid, is a liquid met with in the preparation of ether. The iodide (EtI), hydride (EtH), acetate ($Et\bar{A}$), and other salts are of considerable chemical interest, but not used in medicine.

Absolute or Real Alcohol (C_2H_5HO) may be prepared from spirit of wine by removing the water which the latter contains. This is accomplished, partially, by anhydrous carbonate of potassium, and finally and entirely by recently burnt quicklime. In operating on, say, one pint, $1\frac{1}{2}$ oz. of dried carbonate of potassium is placed in a bottle that can be well closed, and frequently shaken during two days with the spirit. Meanwhile about half a pound of good quicklime, if not already at hand, is made from 10 or 12 oz. of slaked lime by heating to redness in a covered crucible for half an hour. The spirit having been decanted from the denser aqueous solution of carbonate of potassium and placed in a quart flask, retort, or tin can, the lime, as soon as cold, is added, and the whole occasionally shaken during a day. The vessel is now placed in a saucepan or other bath containing water, quickly connected with a condenser (in the case of the flask or can by a bent tube and cork previously prepared; for absolute alcohol must not be exposed to air, or water in the form of moisture will be rapidly reabsorbed), and heat applied to the bath. Rejecting the first ounce or ounce and a half, as likely to contain traces of moisture absorbed from the air or apparatus, continue distillation until nothing more passes over, the water in the bath being kept just below the boiling-point (about

200° F.). These details are those of the British Pharmacopœia. Sp. gr. '7938. Boiling-point 173·6 F.

Tests.—There are no specific tests for alcohol when mixed with complex matters. It is, however, easily isolated and concentrated by fractional distillation, and is then recognisable by conjoint physical and chemical characters. Thus its odour and taste are characteristic; it is lighter than water, volatile, colourless, and, when tolerably strong, inflammable, burning with an almost non-luminous flame; it readily yields aldehyd (*see below*) and acetic ether (*vide Index*) each of which has a characteristic odour; lastly, in presence of hot acid, alcohol reduces red chromate of potassium to a green salt of chromium.

According to Lieben, 1 of alcohol in 2000 of water can be detected by adding to some of the warmed liquid a little iodine, a few drops of solution of soda, again warming gently, and setting aside for a time; a yellowish crystalline deposit of *iodoform* (CHI_3) is obtained. Under the microscope the latter presents the appearance of hexagonal plates or six-rayed and other varieties of stellate crystals.



Other alcohols, aldehyds, gum, turpentine, sugar, and several other substances give a similar reaction.

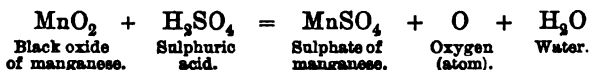
Tests of purity.—Oil or resin is precipitated on diluting spirit of wine with distilled water, giving an opalescent appearance to the mixture. The specific gravity should be 0·838. Fusel oil, aldehyd, and such impurities are detected by nitrate of silver (*vide Index*, "Alcohol, test for purity of"). Water in absolute alcohol may be detected by adding to a small quantity a little highly dried sulphate of copper, which becomes blue ($\text{CuSO}_4, 5\text{H}_2\text{O}$) if water is present, but retains its yellowish-white anhydrous character (CuSO_4) if water be absent.

ALDEHYD ($\text{C}_2\text{H}_4\text{O}$).—Place together, in a capacious test-tube or a flask, about four parts of spirit of wine, six of black oxide of manganese, six of sulphuric acid, and four of water, and gently warm the mixture; aldehyd (*alcohol dehydrogenatus*), a highly volatile liquid, is immediately formed, and its vapour evolved, recognised by its peculiar, somewhat fragrant odour. Adapt a cork and rather long bent tube to the

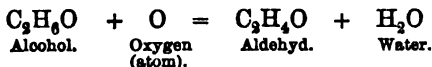
test-tube, and let some of the aldehyd slowly distil over into another test-tube, the condensing-tube being kept as cool as possible. Set the distillate aside for a day or two; the aldehyd will have nearly all disappeared, and acetic acid be found in the tube. Test the exposed liquid by litmus-paper; it will be found to have an acid reaction: make it slightly alkaline by a drop or two of solution of carbonate of sodium, then boil to remove any alcohol and aldehyd present, add sulphuric acid, and notice the characteristic odour of the acetic acid evolved.

These experiments will enable the process of acetification described in connection with acetic acid to be more fully understood. Pure diluted alcohol is not oxidized by exposure to air; but in presence of fermentive matter, or vegetable matter undergoing decay or change, it is oxidized first to aldehyd and then to acetic acid.

In the above process the black oxide of manganese and sulphuric acid furnish nascent oxygen:—



The nascent oxygen then acts on the alcohol, just as the oxygen of the air acts on the alcohol in fermented infusion of malt, beer, or wine, giving aldehyd:—



The aldehyd rapidly, even when pure (more rapidly when impure), absorbs oxygen and yields acetic acid:—



Tests.—Aldehyd heated with solution of potash gives a brownish-yellow resinous mass of peculiar odour. Its aqueous solution reduces salts of silver, giving a mirror-like coating to the sides of a test-tube.

Spirit of French Wine (*Spiritus Vini Gallici*, B. P.) or *Brandy* is a coloured and flavoured variety of alcohol distilled from French wine. Its colour is that of light sherry, and is derived from the cask in which it has been kept, but is com-

monly deepened by the addition of burnt sugar. Its taste is due to the volatile flavouring constituent of the wine, often increased by the addition of artificial essences.

QUESTIONS AND EXERCISES.

751. Write a few sentences on the formation, purification, and concentration of alcohol, and explain the difference between Rectified Spirit, Proof Spirit, and Absolute Alcohol.

752. What quantity of water must be added to one gallon of spirit of wine, 56 degrees over proof, to convert it into proof spirit?

753. To what volume must five pints of spirit of wine of 53 degrees over proof be diluted before it becomes proof spirit?—*Ans.* 7 pints, 13 ounces.

754. State the specific gravity of proof spirit.

755. Show how the formula of alcohol is obtained from its centesimal composition:—Carbon, 52.174; Hydrogen, 13.043; Oxygen, 34.783 = 100.000.

756. Give the formulæ of some of the salts of ethyl.

757. By what processes may pure hydrate of ethyl be obtained?

758. Enumerate the characters of alcohol.

759. Mention a chemical test to distinguish rectified spirit from absolute alcohol.

760. From the formula of aldehyd calculate back its composition in 100 parts.

761. What is the relation of aldehyd to alcohol and to acetic acid?

762. Whence is brandy obtained? and to what are due its colour and flavour?

ETHER, OR OXIDE OF ETHYL.

Formula $C_4H_{10}O$, or $(C_2H_5)_2O$, or Et_2O .

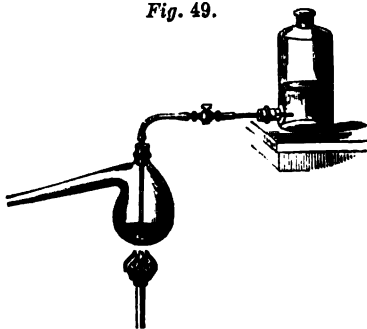
Experimental Process.—Into a capacious test-tube put a small quantity of spirit of wine and about half its bulk of sulphuric acid, mix, and gently warm; the vapour of ether, recognised by its odour, is evolved. Adapt a cork and long bent tube to the test-tube and slowly distil over the ether into another test-tube. Half the original quantity of alcohol now

placed in the generating-tube will again give ether; and this operation may be repeated many times.

On the larger scale, and according to the following official process (*Æther*, B. P.), the addition of alcohol, instead of being intermitting, is continuous, a tube conveying alcohol from a reservoir into the generating-vessel. Mix ten fluid ounces of sulphuric acid with twelve fluid ounces of rectified spirit in a glass retort or flask capable of containing at least two pints, and, not allowing the mixture to cool, connect the retort or flask, by means of a bent glass tube, with a Liebig's condenser, and distil with a heat sufficient to maintain the liquid in brisk ebullition. (If a thermometer also be inserted in the tubulure of the retort or through the cork of the flask, the temperature may be still more carefully regulated—between 284° and 290°F.) As soon as the ethereal fluid begins to pass over, supply fresh spirit in a continuous stream, and in such quantity as to about equal the volume of the fluid which distils. For this purpose use a tube furnished with a stopcock to regulate the

Fig. 49.

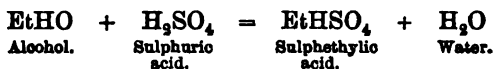
supply, as shown in fig. 49, connecting one end of the tube with a vessel containing the spirit supported above the level of the retort or flask, and passing the other end through the cork of the retort or flask into the liquid. When a total of fifty fluid ounces of spirit has been added, and forty-two fluid ounces of ether have distilled over, the process may be stopped.



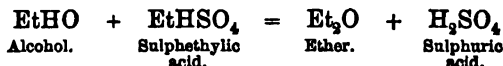
PREPARATION OF ETHER.

To *partially purify* the liquid, dissolve ten ounces of chloride of calcium in thirteen ounces of water, add half an ounce of lime, and agitate the mixture in a bottle with the impure ether. Leave the mixture at rest for ten minutes, pour off the light supernatant fluid, and distil it with a gentle heat until a glass bead of specific gravity 0.735 placed in the receiver begins to float. The ether and spirit retained by the chloride of calcium and by the residue of each rectification may be recovered by distillation and used in a subsequent operation.

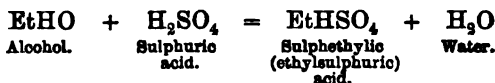
Explanation of Process.—On the addition of sulphuric acid to alcohol in equal volumes, one molecule of each react and give a molecule of sulphethylic acid and one of water:—



More alcohol then gives ether and sulphuric acid by the reaction of one molecule of the alcohol on one of sulphethylic acid:—



The water of the first reaction and the ether of the second distil over, while the sulphuric acid, as fast as liberated, is attacked by alcohol and reconverted into sulphethylic acid,



so that the sulphuric acid originally employed finally remains in the retort in the form of sulphethylic acid. The effect, however, of a small quantity of sulphuric acid in thus converting a large quantity of alcohol into ether is limited, secondary reactions occurring to some extent after a time.

Properties.—Pure ether is gaseous at temperatures above 95° F.; hence the condensing-tubes employed in its distillation must be kept as cool as possible. At all ordinary temperatures it rapidly evaporates, absorbing much heat from the surface on which it is placed. A few drops evaporated consecutively from the back of the hand produce great cold; if blown in the form of spray, the cooling effect is so rapid and intense as to produce local anæsthesia. Its vapour is very heavy, more than twice and a half that of air, and nearly forty times that of hydrogen ($\text{H}_2 = 2$; $\text{C}_4\text{H}_{10}\text{O} = 74$; or as 1 to 37). In a still atmosphere, therefore, it will flow a considerable distance along a table or floor before complete diffusion occurs: the vapour is also highly inflammable; hence the importance of keeping candle and other flames at a distance during manipulations with ether.

Purification.—To imitate the process of partial purification above described add to the small quantity of ether obtained in the foregoing operation a strong solution of chloride of calcium

and a little slaked lime; the latter absorbs any sulphurous acid that may have been produced by secondary decompositions, while the former absorbs water; on shaking the mixture and then setting aside for a minute or two, the ether will be found floating on the surface of the solution of chloride of calcium.

This ether, redistilled until the distillate has a sp. gr. not higher than 0.735, and boiling-point not higher than 105° F., is the ether of the British Pharmacopœia. It still contains about 8 per cent. of alcohol. The latter may be removed by well shaking the ether with half of its bulk of water, setting aside, separating the floating ether and again shaking it with water; alcohol is thus washed out. This washed ether containing water (for water and ether are to some extent soluble the one in the other; fifty measures of ether agitated with an equal volume of water are reduced to forty-five measures) is next placed in a retort with solid chloride of calcium and a little caustic lime, and once more distilled; pure dry ether (*Æther Purus*, B. P.) results. Sp. gr. not exceeding 0.720.

Spiritus Ætheris, B. P., is a mixture of common ether (*Æther*, B. P.) with twice its bulk of rectified spirit.

NITROUS ETHER, OR NITRITE OF ETHYL.

Formula $C_2H_5NO_2$ or $EtNO_2$.

Process.—To a third of a test-tubeful of rectified spirit add about a tenth of its bulk of sulphuric acid, rather more of nitric acid, and some copper wire or turnings, and warm the mixture; as soon as ebullition commences, the vapour of nitrous ether (with other substances) is evolved, recognised by its odour. A long bent tube, kept cool, may be adapted by a perforated cork to the test-tube, and thus a little of the product be condensed and collected.

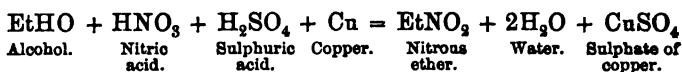
*The above process conducted on a larger scale, with definite quantities of materials, temperature regulated by a thermometer, and a well-cooled condenser, etc. (see page 138), is the official (Redwood's) process for the preparation of a concentrated solution of nitrous ether, etc., in spirit; diluted with nearly three times its bulk of rectified spirit it forms one variety of the "sweet spirit of nitre" (*Spiritus Ætheris Nitrosi*, B. P.) of pharmacy.*

“Take of

Nitric Acid	3 fluid ounces,
Sulphuric Acid	2 fluid ounces,
Copper, in fine wire (about No. 25).	2 ounces,
Rectified Spirit	a sufficiency.

“To one pint of the spirit add gradually the sulphuric acid, stirring them together; then add, in the same way, two and a half fluid ounces of the nitric acid. Put the mixture into a retort or other suitable apparatus, into which the copper has been introduced, and to which a thermometer is fitted. Attach now an efficient condenser, and, applying a gentle heat, let the spirit distil at a temperature commencing at 170°F. and rising to 175°, but not exceeding 180°, until twelve fluid ounces have passed over and been collected in a bottle kept cool, if necessary, with ice-cold water; then withdraw the heat, and, having allowed the contents of the retort to cool, introduce the remaining half-ounce of nitric acid, and resume the distillation as before, until the distilled product has been increased to fifteen fluid ounces. Mix this with two pints of the rectified spirit, or as much as will make the product correspond to the tests of specific gravity and percentage of liquid separated by chloride of calcium (*vide infra*). Preserve it in well-closed vessels.”

Disregarding the other products (including much aldehyd), the following equation probably represents decompositions that occur in the operation. The main point in the reaction is the reduction of the nitric to the nitrous radical by the indirect agency of the copper.



Properties.—Spirit of Nitrous Ether “is transparent and nearly colourless, with a very slight tinge of yellow, mobile, inflammable, of a peculiar penetrating apple-like odour, and sweetish cooling sharp taste. Specific gravity (B. P.) 0.845. It effervesces feebly, or not at all, when shaken with a little bicarbonate of soda” (showing absence of appreciable quantities of free nitrous, acetic, or other acids). The aldehyd in it may be detected by the potash test (*see* p. 507). The great tendency of aldehyd to become converted into acetic acid by the absorption of oxygen from the air renders Spirit of

Nitrous Ether unstable, and pharmacists are obliged to neutralize such acid, generally by bicarbonate of potassium, before adding it to medicines containing iodine, etc.

A second variety of "sweet spirit of nitre" still largely sold in Great Britain, is made from spirit of wine and nitric acid, as ordered in the London Pharmacopœias, except that the distillation is continued until the product contains the maximum of nitrous ether, etc. (i.e. has a sp. gr. of 0.850).

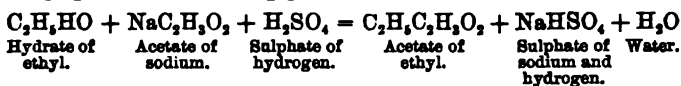
Test.—The nitrous radical may be detected by adding sulphate of iron and sulphuric acid to some of the spirit of nitrous ether, a brown or black compound being produced, already explained in connection with nitric acid.

Official test of strength.—To some of the official Spirit of Nitrous Ether add twice its bulk of a saturated solution of chloride of calcium and mix the liquids; on setting aside, an ethereal fluid will rise to the surface if the spirit has been recently made. If the spirit of nitrous ether be of official strength, not less than 2 per cent. of its volume will thus separate, indicating, it is stated, the presence of 10 per cent. of the ethereal fluid, 8 per cent. being said to be still in solution. A graduated tube is obviously most convenient for this experiment. The "ethereal fluid" is a mixture of aldehyd, nitrous ether, alcohol and other bodies.

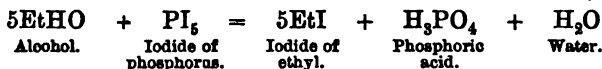
Acetic Ether, or Acetate of Ethyl.

To a little dried acetate of sodium, in a test-tube, add a small quantity of rectified spirit of wine and some sulphuric acid, and, adapting a long bent tube in the usual manner, heat the test-tube and so distil over acetic ether—which may be collected in another test-tube kept cool by partial immersion in cold water.

The official proportions (*Æther Aceticus*, B. P.) are 8 parts of dried acetate of sodium, 5 of spirit, and 10 of acid. It is purified from any water by shaking in a bottle with fused chloride of calcium, and, after twenty-four hours, rectifying. "Sp. gr. 0.910. Boiling point 166° F."



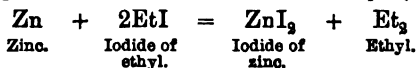
IODIDE OF ETHYL (EtI) may be prepared by mixing two or three parts of phosphorus with 100 of *absolute* alcohol and then dropping in iodine gradually until about 200 parts have been added.



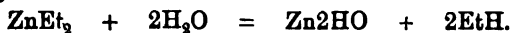
The reaction at first proceeds rapidly, and is complete after the mixture has been set aside for a few hours. The iodide of ethyl may then be isolated by careful distillation, freed from any excess of iodine by washing with a very small quantity of solution of potash or soda, washed with water, dried over chloride of calcium, and again distilled. It should be kept in a dark place, as light favours decomposition and liberation of iodine.

Bromide of ethyl ($\text{C}_2\text{H}_5\text{Br}$) or *hydrobromic ether* may be similarly prepared. For its preparation on a large scale De Vrij's method is preferable, $\text{EtHSO}_4 + \text{KBr} = \text{EtBr} + \text{KHSO}_4$ (*see Pharm. Journ.*, Feb. 15, 1879), or the same method as modified by Greene (*P. J.*, July 12, 1879); by Remington (*P. J.*, May 29, 1880); or by Wolff (*P. J.*, July 3, 1880).

ETHYL.—This gaseous radical, $(\text{C}_2\text{H}_5)_2$ or Et_2 , is obtained on digesting together at about 250°F. , in a strong sealed tube, dry freshly granulated zinc with iodide of ethyl (Frankland).



On cautiously opening the tube the ethyl escapes, and may be ignited or collected over water. There remains with the iodide of zinc a body termed by Frankland *zinc-ethyl* (ZnEt_2); it is a spontaneously inflammable liquid, but may easily be distilled and otherwise manipulated if a few simple precautions be observed. If water be allowed to flow down the tube, the solid compound of iodide of zinc and zinc-ethyl will be decomposed, a gas, *hydride of ethyl* (EtH), resulting, which also may be inflamed or collected over water:—



QUESTIONS AND EXERCISES.

763. Describe the official process for the preparation of Ether, giving equations.

764. Offer a physical explanation of the mode of producing local anæsthesia.

765. How is commercial ether purified?

766. Explain the official process for the preparation of Spirit of Nitrous Ether.

767. Give the properties of spirit of nitrous ether.

768. By what official method is the strength of spirit of nitrous ether to be estimated?

769. How is iodide of ethyl made?

770. Adduce evidence of the existence of ethyl.

OTHER ALCOHOL RADICALS AND THEIR SALTS.

What has been stated concerning the chemistry of ethyl and its compounds may be applied to other radicals known to exist, some of the compounds of each of which are of common occurrence. These basylous radicals are closely related to each other, to hydrogen, and to the metals. Their formulæ (in the combined state) may be built up by successive additions of CH_2 , thus:—

Hydrogen	H
Methyl	CH_3 , or Me
Ethyl	C_2H_5 , or Et
Propyl (or Trityl)	C_3H_7 , or Pr
Butyl (or Tetryl)	C_4H_9 , or Bu
Amyl	C_5H_{11} , or Ay
Caproyl (or Hexyl)	C_6H_{13} , or Cp

The above list is an illustration of an *homologous* series (from $\delta\mu\delta\varsigma$, *homos*, the same, and $\lambda\acute{o}\gamma\omicron\varsigma$, *logos*, proportion) of compounds. It will be observed that the relation of the number of hydrogen atoms to carbon is twice as many with one added; hence the series is often termed the $\text{C}_n\text{H}_{2n+1}$ series (n = any number). The oxides of these radicals are known as *ethers*, their hydrates *alcohols*, their compounds with the acetic and similar acidulous radicals *ethereal salts*. Every alcohol furnishes a body corresponding to the aldehyd of spirit of wine, the class being termed *aldehyds*; each also yields an

acid corresponding with acetic acid. Any one of these classes constitutes an homologous series. Or, taking the hydride, oxide, hydrate, acid, of any single radical, we get a *heterologous* (*irepos, heteros*, another) series of compounds. Hydride of methyl (MeH or CH_3H) is ordinary *marsh-gas, fire-damp*, or *light carburetted hydrogen*; it is a diluent or non-luminiferous constituent of ordinary coal-gas to the extent of 30 to 40 per cent.*; *formic acid*, the acid of the methyl series; *butyric acid*, the acid of the butyl series; *sulphocyanate of butyl*, the essential oil of horseradish; *valerianic acid*, the acid of the amyl series.

Homologous and Heterologous Series of the $\text{C}_n\text{H}_{2n+1}$ Radicals.

Radicals. (Free).	Hydrides.	Oxides (or ethers).	Hydrates (or alcohols).	Aldehyds.	Acids.
$(\text{CH}_3)_2$	CH_3H	$(\text{CH}_3)_2\text{O}$	CH_3HO	$\text{CH}_3\text{O} ?$	CH_3O_2
$(\text{C}_2\text{H}_5)_2$	$\text{C}_2\text{H}_5\text{H}$	$(\text{C}_2\text{H}_5)_2\text{O}$	$\text{C}_2\text{H}_5\text{HO}$	$\text{C}_2\text{H}_5\text{O}$	$\text{C}_2\text{H}_5\text{O}_2$
$(\text{C}_3\text{H}_7)_2$	$\text{C}_3\text{H}_7\text{H}$	$(\text{C}_3\text{H}_7)_2\text{O}$	$\text{C}_3\text{H}_7\text{HO}$	$\text{C}_3\text{H}_7\text{O}$	$\text{C}_3\text{H}_7\text{O}_2$
$(\text{C}_4\text{H}_9)_2$	$\text{C}_4\text{H}_9\text{H}$	$(\text{C}_4\text{H}_9)_2\text{O}$	$\text{C}_4\text{H}_9\text{HO}$	$\text{C}_4\text{H}_9\text{O}$	$\text{C}_4\text{H}_9\text{O}_2$
$(\text{C}_5\text{H}_{11})_2$	$\text{C}_5\text{H}_{11}\text{H}$	$(\text{C}_5\text{H}_{11})_2\text{O}$	$\text{C}_5\text{H}_{11}\text{HO}$	$\text{C}_5\text{H}_{11}\text{O}$	$\text{C}_5\text{H}_{11}\text{O}_2$
etc.,	etc.	etc.	etc.	etc.	etc.

QUESTIONS AND EXERCISES.

771. Mention several radicals homologous with ethyl, and give their formulæ.

772. Define ethers, hydrides, alcohols, ethereal salts, aldehyds.

773. What is the difference between homologous and heterologous series?

774. Give the systematic name of *fire-damp*.

* *Coal-gas*.—The other diluents, or vehicles for the conveyance of illuminating vapours, of coal-gas are hydrogen (40 to 50 per cent.) and carbonic oxide (6 to 7 per cent.). The illuminating constituents are olefant gas (p. 533) and its homologues, existing to the extent of from 5 to 7 per cent. Hydrocarbons, normally fluid but kept in the vaporous condition by the diluents, also contribute materially to the illuminating power of gas. The impurities are nitrogen, air, carbonic acid, bisulphide of carbon CS_2 (a volatile liquid easily made from its elements and rendered almost scentless by digesting with copper turnings), and other badly smelling sulphur compounds. Upwards of fifty distinct chemical substances have been obtained from the solid, liquid, and gaseous products of the destructive distillation of coal.

775. Enumerate the chief constituents of coal-gas. .

776. State the formulæ of formic, butyric, and valerianic acids.

777. Write the formulæ of butyl, its hydride, ether, alcohol, aldehyd, and acid.

METHYLIC ALCOHOL.

METHYLIC ALCOHOL (CH_3HO , or MeHO), WOOD-SPIRIT, or PYROXYLIC SPIRIT, or WOOD-NAPHTHA, is a product of the destructive distillation of wood. Spirit of wine containing 10 per cent. of wood-spirit constitutes ordinary *methylated spirit*, a spirit issued duty free, for the use of manufacturers, the methylic alcohol not interfering with technical applications. From its nauseous taste and odour, however, it cannot take the place of gin, brandy, or other spirit; hence, while industry is benefited, intemperance is discouraged and the revenue not injured.

Detection of Methylic Alcohol in presence of Ethylic Alcohol.—Three or four methods have been proposed for the detection of methylated spirit in various liquids; that open to least objection is by J. T. Miller. For the application of the test to tinctures and similar spirituous mixtures, some of the spirit is first separated by distilling off a drachm or so from about half an ounce of the liquid placed in a small flask or test-tube having a long bent tube attached. Into a similar apparatus put 30 grains of powdered red chromate of potassium, half an ounce of water, 25 minims of strong sulphuric acid, and 30 or 40 minims of the spirit to be tested. Set the mixture aside for a quarter of an hour and then distil nearly half a fluid ounce. Place the distillate in a small dish, add a very slight excess of carbonate of sodium, boil down to about a quarter of an ounce, add enough acetic acid to impart a distinct but feeble acid reaction, pour the liquid into a test-tube, add a grain of nitrate of silver dissolved in about 30 drops of water, and heat gently for a couple of minutes. If the liquid then merely darkens a little, but continues quite translucent, the spirit is free from methylic alcohol; but if a copious precipitate of dark-brown or black metallic silver separates, and

the tube, after being rinsed out and filled with clean water, has a distinct film of silver, which appears brown by transmitted light (best seen by holding it against white paper), the spirit is methylated.

Explanation.—This test depends for its action on the reducing-powers of formic acid. In the above operation the ethylic alcohol becomes oxidized to acetic acid (the natural acid of the ethyl series), which does not reduce silver salts, a minute quantity only of formic acid being produced, while the methylic alcohol yields formic acid (the natural acid of the methyl series) in a comparatively large quantity. Aldehyd, which is also a reducing agent, is simultaneously produced, but removed in the subsequent ebullition with carbonate of sodium.

Methylated Sweet Spirit of Nitre.—The preparation of spirit of nitrous ether from methylated spirit is illegal in Great Britain, and, probably, is very rarely practised. For the detection of methylic alcohol in this liquid, Mr. Miller suggests the following modification of the above process.

Shake about an ounce of the sample with 20 or 30 grains of anhydrous carbonate of potassium, and, if needful, add fresh portions of the salt until it ceases to be dissolved, then pour off the supernatant spirit. This serves to neutralize acid and to remove water, of which an abnormal quantity may be present. Introduce half a fluid ounce of the spirit into a small flask; add 150 grains of anhydrous chloride of calcium in powder, and stir well together; then, having connected the flask with a condenser, place it in a bath of boiling water, and distil a fluid drachm and a half, or continue the distillation until scarcely anything more comes over. The operation is rather slow, but needs little attention, and should be done thoroughly. The distillate contains nearly the whole of the nitrous ether and other interfering substances, while in the retort there remains a non-volatile compound of chloride of calcium and methylic alcohol, if the latter be present. Now add to the contents of the flask a fluid drachm of water, which decomposes the compound just referred to, and draw

over the half-drachm of spirit required for testing. Add it to the usual oxidizing solution composed of 30 grains of red chromate of potassium, 25 minims of strong sulphuric acid, and half an ounce of water; let the mixture stand a quarter of an hour, then distil half a fluid ounce. Treat the distillate with a slight excess of carbonate of sodium, boil rapidly down to two fluid drachms, and drop in, cautiously, enough acetic acid to impart a faint acid reaction; pour the liquor into a test-tube about three-quarters of an inch in diameter; add two drops of *diluted* acetic acid, B.P., and one grain of nitrate of silver in half a drachm of pure water; apply heat, and boil gently for two minutes. If the spirit is free from methylic alcohol the solution darkens and often assumes transiently a purplish tinge, but continues quite translucent, and the test-tube, after being rinsed out and filled with water, appears clean or nearly so. But if the spirit contains only 1 per cent. of methylic alcohol the liquid turns first brown, then almost black and opaque, and a film of silver, which is brown by transmitted light, is deposited on the tube. When the sample is methylated to the extent of 3 or 4 per cent., the film is sufficiently thick to form a brilliant mirror. To insure accuracy, the experiment should be performed by daylight.

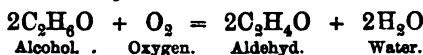
CHLOROFORM.

Formula CHCl_3 .

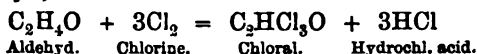
Process.—Should the necessary appliances be at hand, a small quantity of this liquid may easily be prepared by the official process. One fluid ounce and a half of spirit and 24 of water are placed in a retort or flask of at least a quart capacity; 8 oz. of chlorinated lime and 4 of slaked lime are added, the vessel connected with a condenser, and the mixture heated until distillation commences, the source of heat then being withdrawn. The condensed liquid should fall into a small flask containing water, at the bottom of which about a drachm of chloroform will slowly collect.

Explanation of the Process.—The hypochlorite of calcium believed to be present in the chlorinated lime (*see* the re-

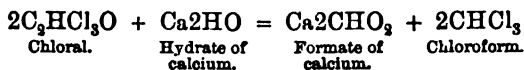
marks in connection with the latter, p. 122) readily yields up oxygen and chlorine to organic substances, the calcium being liberated as hydrate, $4(\text{Ca}_2\text{ClO}) + 4\text{H}_2\text{O} = 4(\text{Ca}_2\text{HO}) + 2\text{O}_2 + 4\text{Cl}_2$. The alcohol used in making chloroform is thus probably first reduced to aldehyd:—



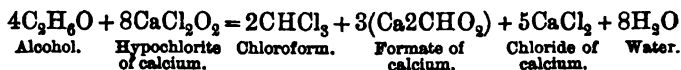
The action of chlorine on aldehyd then probably gives chloral (*chlor-aldehyd*):—



The hydrochloric acid being at once neutralized by some of the liberated hydrate of calcium to form chloride of calcium and water, more freed hydrate of calcium and chloral give formate of calcium and chloroform.



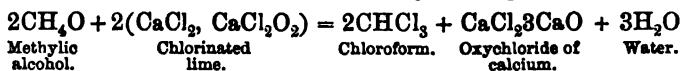
Or, neglecting the probable steps in the process, and regarding only the materials and the products, 4 molecules of alcohol and 8 of hypochlorite of calcium give 2 of chloroform, 3 of formate of calcium, 5 of chloride of calcium, and 8 of water, thus:—



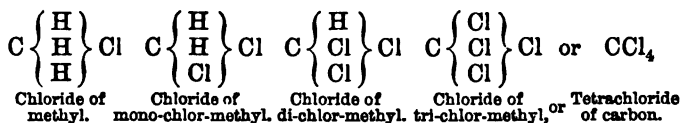
The hydrate of calcium placed in the generating-vessels is not essential, but is useful in preventing secondary decompositions, the hydrate of calcium obtainable from the reaction being insufficient for this purpose.

Constitution.—Chloroform is sometimes considered to be the chloride of a trivalent radical *formyl* (CH), the first member of a series $\text{C}_n\text{H}_{2n-1}$. Glycerine is the hydrate of another member—*glyceryl*, C_3H_5 (p. 533).

Chloroform may also be regarded as the chloride of dichlor-methyl; it may be formed from methylic compounds, thus:—



Chlorine converts it into tetrachloride of carbon, completing a series of *substitution products* of chloride of methyl.



The chloride of mono-chlor-methyl, under the name of di-chloride of methylene, has been used as an anæsthetic. It may be obtained by the action of nascent hydrogen on chloroform.

Chloroform is purified by shaking it with water and then with pure sulphuric acid (containing no trace of nitric acid), which chars and removes hydrocarbons, etc., but does not affect chloroform. It is freed from any trace of acid by agitation with lime, and from moisture by solid chloride of calcium.

Properties.—The sp. gr. of chloroform is 1.500. It is liable to slowly decompose when exposed to air and light. To render it stable a minute amount of alcohol is necessary; hence the specific gravity of medicinal chloroform is about 1.497. It readily and entirely volatilizes at common temperatures, having, to the last drop, its pleasant characteristic odour. It has a sweetish taste, is limpid, colourless, soluble in alcohol and ether, and slightly in water. It burns with a sluggish green smoky flame. It should be neutral to test paper, indicating absence of acid; give no precipitate with solution of nitrate of silver, indicating absence of ordinary chlorides; remain colourless when heated with potash, indicating absence of aldehyd; and give no colour to any sulphuric acid with which it may be shaken, even after the mixture has been set aside for half an hour, indicating absence of hydrocarbons, etc. Alcohol may be detected by Lieben's test (p. 506), or by shaking with a little of the dye termed "Hofmann's violet," which gives the chloroform a purple tint if alcohol be present, but affords no colour with pure chloroform.

Aqua Chloroformi, B. P., the official Chloroform Water, is made by shaking 1 fluid drachm of chloroform with 25 ounces of distilled water till dissolved.

IODOFORM (CHI_3), analogous in constitution to chloroform, the iodine occupying the place of the chlorine, is made by mixing in a retort 2 parts of carbonate of potassium, 2 parts of iodine, 1 part of alcohol, and 5 of water, heating till colourless, and then pouring into a beaker and allowing to settle. The iodoform is deposited in yellow scales, which are collected on a filter, washed thoroughly with water, and dried between filtering-paper.

Iodoform appears in the shape of yellow, shining, six-sided scales. It is volatile at ordinary temperatures. Almost insoluble in water, but more so in alcohol.

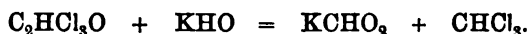
CHLORAL AND CHLORAL HYDRATE.

Process.—Pass a rapid stream of dry chlorine into pure absolute alcohol so long as absorption occurs. During the first hour or two the alcohol must be kept cool, afterwards gradually warmed till ultimately the boiling-point is reached. The preparation of a considerable quantity occupies several days. The crude product is mixed with three times its volume of oil of vitriol and distilled, again mixed with a similar quantity of oil of vitriol and again distilled, and finally rectified from quicklime.

The formation of chloral would at first sight seem to be due to the production from the alcohol (C_2H_5O) of aldehyd (C_2H_4O), through the removal of hydrogen by the chlorine, and the substitution of chlorine for hydrogen in the aldehyd (C_2H_4O) with formation of *chlor-aldehyd* or *chloral* (C_2HCl_3O). But the reactions are far more complicated, being somewhat as follows. Aldehyd and hydrochloric acid are first formed; these with some of the alcohol give monochlorinated ether, $C_2H_5 \left. \begin{array}{l} C_2H_4Cl \\ C_2H_4Cl \end{array} \right\} O$, which with more chlorine yields tetrachlorinated ether $C_2H_5 \left. \begin{array}{l} C_2H_5 \\ C_2HCl_4 \end{array} \right\} O$, and this in presence of water furnishes chloral, some alcohol, and hydrochloric acid (Wurtz and Vogt).

Properties.—The formula of chloral is C_2HCl_3O . It is a colourless liquid, of oily consistence. Sp. gr. 1.502. Boiling point $201.2^\circ F$. Its vapour has a penetrating smell, and is somewhat irritating to the eyes. Mixed with water heat is disengaged, and solid white, crystallizable, *hydrous chloral*, or what is more generally though somewhat irregularly termed *chloral hydrate* (C_2HCl_3O, H_2O) or *Hydrate of Chloral* (*Chloral Hydras*, B. P.), results. The latter fuses at 110.8° and boils at $203^\circ F$. It sublimes as a white crystalline powder. Both chloral and chloral hydrate are soluble in water, alcohol, ether, and oils. Oils and fats are also soluble in chloral hydrate. The aqueous solution should be neutral, and give no reaction with nitrate of silver. Chloral hydrate is said sometimes to

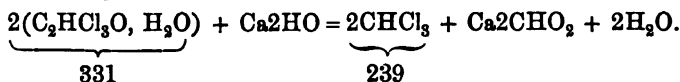
undergo a spontaneous change into an opaque white isomeric modification, insoluble in water, alcohol, or ether, but convertible by prolonged contact with water or by distillation into the ordinary condition. By action of weak alkalies chloral first yields formate of the alkali-metal and chloroform:—



Chloral, or rather strong aqueous solution of chloral hydrate (3 in 4), injected beneath the skin yields nascent chloroform by action of the alkali of the blood, and produces narcotic effects (Liebreich, Personne). Chloroform itself admits of similar hypodermic use (Richardson). If administered by the stomach, thirty to eighty grains of solid hydrate are required. The final products of the reaction of the chloroform and blood are formate and chloride of sodium. A spirituous solution of potash effects the same transformation.



Solution of ammonia and moist hydrate of calcium, as well as weak solutions of fixed alkalies, convert hydrate of chloral into formate of the metal and chloroform. The reaction with the slaked lime being especially definite and complete (Wood), it may be employed in ascertaining the richness of a sample of commercial chloral hydrate in chemically pure chloral hydrate.



From the foregoing equation and molecular weights it is obvious that 100 grains of hydrate of chloral, if quite dry, will yield by distillation with 30 grains of slaked lime and an ounce of distilled water (in a small flask and long bent tube kept cool by moistened paper) 72·2 grains of chloroform by weight, or (the sp. gr. of chloroform being taken at 1·497) 47·56 grains by measure, or about 52 minims. 100 grains of the official hydrate of chloral "should yield not less than 70 grains of chloroform."

Pure Chloral Hydrate.—Liebreich, who first proposed the use of chloral hydrate, gives the following as the characteristics of a pure article:—Colourless, transparent crystals. Does not decompose by the action of the atmosphere, does not leave oily spots when pressed between blotting-paper, affects neither cork nor paper. Smells agreeably aromatic, but a little pungent when heated. Taste bitter astringent, slightly caustic. Seems to melt on rubbing between the fingers. Dissolves in water like candy without first forming oily drops; and the solution is neutral or faintly acid to test-paper. Dissolves in bisulphide of carbon, petroleum, ether, water, alcohol, oil of turpentine, etc. Its solution in chloroform gives no colour when shaken with sulphuric acid. Boiling-point 203° to 205° F., and volatilizes without residue. Distilled with sulphuric acid, the chloral should pass over at 205° to 207° F. Melting-point 133° to 136° F., again solidifying at about 120° . Gives no chlorine reaction on treating the solution in water (acidulated by nitric acid) with nitrate of silver.

Impure Chloral Hydrate.—Yellowish, cloudy. Decomposes; leaves spots by pressing between blotting-paper; decomposes corks and paper of the packing. Smells pungent and irritating; on opening the bottles is sticky and often emits fumes. Taste strongly caustic. With water forms oily drops or is partially insoluble. Boils at a higher temperature. On treating it with sulphuric acid turns brown, with formation of hydrochloric acid. Gives chlorine reaction on treating the solution in water (acidulated by nitric acid) with nitrate of silver.

Alcoholates of chloral are obtained on combining alcohols with chloral.

Bromal (C_2HBr_3O), *hydrate of bromal* (C_2HBr_3O, H_2O), and *alcoholates of bromal* are produced when bromine instead of chlorine attacks alcohol. *Iodal* (C_2HI_3O) also exists.

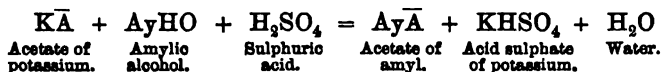
Butyl chloral ($C_4H_5Cl_3O$) originally, but erroneously, termed *croton chloral*, is a product of the action of chlorine on aldehyd. Its name expresses its constitution; it is *chlorinated butyric aldehyd*—ordinary chloral being chlorinated vinic aldehyd.

AMYLIC ALCOHOL.

AMYLIC ALCOHOL (*Alcohol Amylicum*, B. P.) ($C_6H_{11}HO$, or $AyHO$) is a constant accompaniment of ethylic or common alcohol (C_2H_5HO , or $EtHO$) when the latter is prepared from sugar which has been derived from starch; hence the name, from *amylum*, starch. The sugar of potato-starch yields a considerable quantity; hence the alcohol is often called *potato-oil*. It is also termed *fousel-oil*, or *fusel-oil* (from $\phi\acute{u}\omega$, *phuo*, I produce), in allusion to the circumstance that the supposed oil is not simply educed from a substance already containing it, as is usually the case with oils, but is actually produced during the operation. It was described as oil probably because it resembled oil in not readily mixing with water; but it is soluble to some extent in water, and is a true spirit, homologous with spirit of wine. It often contains variable proportions of propylic, butylic, and caproylic alcohols. (See also VALERIANIC ACID.)

Amylic alcohol is "a colourless liquid, with a penetrating and oppressive odour and a burning taste. When pure its specific gravity is .818; boiling-point $270^{\circ}F$. Sparingly soluble in water, but soluble in all proportions in alcohol, ether, and essential oils. Exposed to the air in contact with platinum-black, it is slowly oxidized, yielding valerianic acid." Two allotropic varieties of amylic alcohol exist, one dextro-rotating a polarized ray. The amylic alcohol of trade probably contains both varieties.

ACETATE OF AMYL ($C_6H_{11}C_2H_3O_2$, or $Ay\bar{A}$).—To a small quantity of amylic alcohol in a test-tube add some acetate of potassium and a little sulphuric acid, and warm the mixture; the vapour of acetate of amyl is evolved, recognised by its odour, which is that of the jargonelle pear. If a condensing-tube be attached, the essence may be distilled over, washed by agitation with water to free it from alcohol, and separated by a pipette.

*Fruit-Essences.*

Acetate of amyl, prepared with the proper equivalent proportions of constituents as indicated by the above equation, is

largely manufactured for use as a flavouring agent by confectioners. Valerianate of amyl ($C_6H_{11}C_5H_9O_2$) is similarly used under the name of apple-oil. Butyrate of ethyl ($C_2H_5C_4H_7O_2$) closely resembles the odour and flavour of the pine-apple; cœnanthylate of ethyl ($C_2H_5C_7H_{13}O_2$) recalls greengage; pelargonate of ethyl ($C_2H_5C_9H_{17}O_2$) quince; suberate of ethyl ($Et_2C_8H_{13}O_4$) mulberry; sebacate of ethyl ($Et_2C_{10}H_{16}O_4$) melon. Hydride of salicyl ($C_7H_5O_2H$), or salicylous acid, is the essential oil of meadow-sweet (*Spiræa ulmaria*), and may be prepared artificially by the oxidation of salicin (*vide* page 499). Acid salicylate of methyl ($CH_3HC_7H_4O_3$), or gaultheric acid, forms the chief part of the essential oil of winter-green (*Gaultheria procumbens*, the fresh leaves of which yield about 0.7 per cent. of oil), and may also be prepared artificially from salicin. Salicylic acid ($C_7H_5O_3$ or $H_2C_7H_4O_3$) can easily be obtained from the salicylate of methyl, but more cheaply from carbolic acid (*vide* page 530).

By mixing ethereal salts with each other and with essential oils in various proportions, the odour and flavour of nearly every fruit may be fairly imitated. (For a set of formulæ of fruit essences see *Pharmaceutical Journal*, May 17, 1879.)

Nitrite of Amyl.

NITRITE OF AMYL (*Amyl Nitris*, B. P.) ($C_5H_{11}NO_2$).—For the preparation of this substance Maisch recommends a modification of Balard's process. Amylic alcohol is first rectified until pure; that is, until its boiling-point is about $270^\circ F$. This alcohol, with about an equal bulk of nitric acid, is introduced into a capacious glass retort, and a moderate heat is applied and very gradually increased. As soon as the mixture approaches boiling, the fire is removed, and the reaction allowed to continue. If the application of the heat has been too rapid or too long continued, considerable frothing occurs, and the contents of the retort are apt to foam over. With a moderate and slowly increased heat, the reaction is less violent and the temperature rises gradually after the removal of the fire and the beginning of boiling. As soon as the thermometer, inserted in the tubulus, rises above $212^\circ F$, the receiver is changed, the distillate now becoming more and more mixed with ethyl-amylic ether and nitrate of amyl, readily perceived by the change in odour.

The distillate obtained below $212^\circ F$. is agitated with an aqueous solution of hydrate or carbonate of potassium to re-

move free acids, and, after separation, the oily liquid is introduced into a clean retort and again slowly heated. The first portion coming over contains amylic aldehyd. When the very slowly increased heat has risen to 205° F., the receiver is again changed and the distillate now collected as nitrate of amyl, until the thermometer reaches 212° F., when the distillation is stopped.

Nitrite of amyl is a yellowish ethereal liquid; sp. gr. of liquid 0.877, of vapour 4.03; boiling-point 205° F.; soluble in spirit of wine, insoluble in water; converted by fused caustic potash into valerianate of potassium.

QUESTIONS AND EXERCISES.

778. Name the source of methylic alcohol.

779. What is "methylated spirit"?

780. Describe the method by which methylated spirit is detected in a tincture.

781. In what relation does formic acid stand to methylic alcohol?

782. How would you proceed to ascertain whether or not a specimen of sweet spirit of nitre had been made from methylated spirit?

783. Give details of the production of chloroform from alcohol, tracing the various steps by equations.

784. Is chloroform an ethylic compound? What is its probable constitution?

785. How is chloroform purified?

786. State the characters of pure chloroform.

787. Describe the reactions that occur in the manufacture of chloral and chloral hydrate.

788. What is the nature of the action of alkalies on chloral hydrate?

789. Mention the characters of pure and impure chloral hydrate.

790. Whence is amylic alcohol obtained?

791. Has valerianic acid any chemical relation to amylic alcohol?

792. Mention the systematic names of several artificial fruit-essences.

793. What is the formula of nitrite of amyl? and how is it prepared?

SALTS AND DERIVATIVES OF RADICALS OF OTHER SERIES THAN THE C_nH_{2n+1} .

What has been stated regarding radicals having the general formula C_nH_{2n+1} and their salts, may be applied to the radicals of other series.

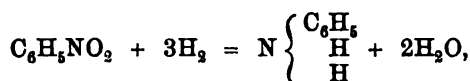
The series C_nH_{2n-7} includes *phenyl* (C_6H_5), the hydride of which (C_6H_5H , or PhH) is common *benzol* or *benzene* (B. P.), a colourless volatile liquid obtained from coal-tar, and a body which in constitution is the type of a very large class of aromatic organic bodies. *Toluol* (C_7H_8), or methyl-benzol, is one of the products of the distillation of balsam of Peru. The next homologue, *xylol* (C_8H_{10}), or dimethyl-benzol, is contained in crude wood-spirit and in coal-naphtha: it is a colourless fluid, having an odour faintly resembling that of benzol; boiling-point $282^\circ F.$; sp. gr. $\cdot 866$. Benzol is a powerful solvent of grease, and under the name of *Benzine Collas* was introduced by M. Collas, in 1848, for cleansing stuffs.* By the action of strong nitric acid, benzol yields *nitrobenzol* [$C_6H_5(NO_2)H$], a liquid termed, from its odour, *artificial oil of bitter almonds*, or *essence of mirbane*. Its specific gravity is 1.20 to 1.29. The odour of this essence, however, is not exactly that of essential oil of almonds, and its composition is very different; so that it is not truly an artificial volatile oil. The natural oil has a sp. gr. of 1.04 to 1.07, and is a hydride of the negative radical† benzoyl (C_7H_5OH), a radical derived from the next higher homologue of phenyl by displacement of hydrogen by oxygen. Nitrobenzol yields aniline (*vide infra*), oil of bitter almonds does not (p. 492). On this reaction the test for nitrobenzol is founded (p. 530), *Dinitrobenzol* [$C_6H_3(NO_2)_2H$] may also be obtained. The hydrate of phenyl (C_6H_5HO), or *phenic alcohol*, or *phenol*, is the *phenic acid* or *carbolic acid* ($H_2C_6H_4O$) of commerce (*Acidum Carbolicum*, B. P.), a colourless crystalline substance obtained from coal-tar oil by fractional distillation and subsequent purification. A crystalline, hydrous, acid ($H_2C_6H_4O, H_2O$) may also be obtained. At temperatures above $95^\circ F.$

* The light products of petroleum are sometimes termed *benzin*, *benzine*, or *benzoline*. They require five or six times their bulk of alcohol for solution, while benzol dissolves in less than its own bulk. Sp. gr. of benzol, about $\cdot 9$; of benzoline, about $\cdot 7$.

† Negative basylous organic radicals are derived from the ordinary or positive basylous organic radicals by displacement of two atoms of hydrogen in the latter radicals by one atom of oxygen.

ordinary carbolic acid is an oily liquid. It is only slightly soluble in water, but readily dissolved by alcohol, ether, and glycerine (*Glycerinum Acidi Carbolici*, B. P.). In odour, taste, and solubility (and in appearance when liquefied by heat or by the addition of 5 per cent. of water) it resembles creasote, a wood-tar product for which carbolic acid is often substituted. Besides hydrate of phenyl (C_6H_5HO) coal-tar oil contains *cresol*, *hydrate of cresyl*, or *cresylic acid* (C_7H_7HO); while wood-tar oil furnishes *guaiacol* ($C_7H_8O_2$)—also a product of the destructive distillation of guaiacum-resin—and *creasol* ($C_9H_{10}O_2$), or *creasote*. Certain colouring-matters may be obtained by the oxidation of carbolic acid; ammonia or, still better, phenyl-ammonia (aniline) mixed with it, and then a small quantity of solution of a hypochlorite, gives a blue liquid; a similar effect is produced on dipping a chip of deal into carbolic acid (or into creasote), then into hydrochloric acid, and afterwards exposing it to the air. By the following tests carbolic acid may be distinguished from creasote. The former boils only at $370^\circ F.$, while the latter readily dries up at 212° . Carbolic acid does not affect a ray of polarized light; creasote twists it slightly to the right. Carbolic acid is either solid or may be solidified by cooling; creasote is not solidified by the cold produced by a mixture of hydrochloric acid and sulphate of sodium. Creasote from coal (impure or crude carbolic acid) gives a jelly when shaken with collodion; creasote from wood (*Oreasotum* B. P.) is unaffected by collodion (Rust). Coal-creasote is soluble in solution of potash and in the strongest solution of ammonia (Read), wood-creasote scarcely soluble. The coal product is soluble in a large volume of water (about 1 in 80, or 1 in 20 of boiling water), and a neutral solution of ferric chloride strikes a blue colour with the liquid; wood-creasote is less soluble (*Aqua Creasoti*, U. S. P., is said to contain 1 in 129) and not coloured blue by ferric chloride. An alcoholic solution of the coal-oil is coloured brown by ferric chloride, a similar solution of true creasote green. According to Mr. Thomas Morson pure creasote is unaffected when mixed with an equal volume of commercial glycerine, while carbolic acid is miscible in all proportions, and will carry into solution even a considerable quantity of creasote. Carbolic acid is a powerful *antiseptic* (*ἀντι*, *anti*, against, and *σῆνω*, *sēpō*, I putrefy). In large doses it is poisonous, *antidotes* being a mixture of olive-oil and castor-oil, freely administered, or a mixture of slaked lime

with about three times its weight of sugar rubbed together with a little water. Carbolic acid is soluble in oil of vitriol, *sulphocarbolic acid* ($\text{HC}_6\text{H}_5\text{SO}_4$) or *sulphophenic acid* being formed. On diluting and mixing with oxides, hydrates, or carbonates, *sulphocarbonates* are formed. The formula of *sulphocarbonate of sodium* is $\text{NaC}_6\text{H}_5\text{SO}_4$; *sulphocarbonate of zinc* $\text{Zn}(\text{C}_6\text{H}_5\text{SO}_4)_2 \cdot \text{H}_2\text{O}$. Tri-nitro-carbolic acid [$\text{C}_6\text{H}_3(\text{NO}_2)_3\text{O}$] is formed on slowly dropping carbolic acid into fuming nitric acid; it is the yellow dye known as *carbazotic acid* or *picric acid*; most of the picrates are explosive by percussion. Both carbolic acid and benzol are secondary products, obtained in the manufacture of coal-gas; hence, indeed, the word *phenic* and thence *phenyl* (from *φαῖνω*, *phainō*, I light, in allusion to the use of coal-gas). *Aniline*, or *phenylamine*, is a product of the action of nascent hydrogen on nitrobenzol.



Nitrobenzol. Hydrogen.

the substance whence, by oxidation, etc., aniline-red (magenta), -orange, -yellow, -green, -blue, -violet, (mauve), and -black are produced. The process with zinc dust (*vide* Watts's Dict.) answers fairly well. One oz. of nitrobenzol gives a drachm or two of aniline. But action of iron filings and acetic acid on an alcoholic solution answers better, 3 oz. of nitrobenzol giving 1 of aniline. To detect nitrobenzol add alcohol and hydrochloric acid and a little zinc; when action has ceased, liberate aniline by shaking with excess of potash; dissolve out the aniline by agitation with ether; to the decanted ethereal liquid add solution of chloride of lime; a violet colour or tint results.

Salicylic acid ($\text{H}_2\text{C}_7\text{H}_4\text{O}_3$).—The natural acid salicylate of methyl is described on page 526. Artificial salicylic acid is made as follows. Carbolic acid ($\text{H}_2\text{C}_6\text{H}_4\text{O}$), by acquiring the elements of carbonic acid gas (CO_2), is converted into salicylic acid (Kolbe). The carbolic acid is first mixed with caustic soda in molecular proportions and dried. The resulting acid carbonate of sodium ($\text{NaHC}_6\text{H}_4\text{O}$) made hot is saturated with carbonic acid gas, every pair of molecules of the carbonate thus affording one of regenerated carbolic acid, which distils off, and one of normal carbonate of sodium ($\text{Na}_2\text{C}_6\text{H}_4\text{O}$) which, absorbing the gas, becomes normal salicylate of sodium

($\text{Na}_2\text{C}_7\text{H}_4\text{O}_3$). The latter by the action of hydrochloric acid furnishes salicylic acid, which may be purified by recrystallization from alcohol. Salicylic acid, like carbolic acid, is a powerful antiseptic, but is free from the taste and smell of carbolic acid. It is only slightly soluble in water, but readily soluble in aqueous solutions of such alkali-metal salts as borax, phosphate of sodium, or citrate of potassium. A similar antiseptic, *cresotic acid* ($\text{C}_8\text{H}_8\text{O}_3$), is similarly obtained from *cresylic acid* ($\text{C}_7\text{H}_8\text{O}$). Ferric chloride strikes a violet coloration with both salicylic and cresotic acids. Both acids have antipyretic powers. The true salicylates of the alkali-metals, and probably therefore the cresotates, are very feeble antiseptics. Salicylate of sodium is fairly soluble in water. Carbolic acid often containing cresylic acid, commercial salicylic acid may contain cresotic acid. Salicylic acid is soluble in strong sulphuric acid yielding *sulphosalicylic acid* $\text{HC}_7\text{H}_5\text{O}_3\text{SO}_4$.

Aloins.—The aloes of pharmacy (*Aloe Barbadosensis*, B. P., and *Aloe Socotrina*, B. P.) is an evaporated juice, doubtless much altered by the temperature to which it is subjected.

Barbaloin.—This substance, first obtained by T. and H. Smith, occurs in minute crystals in Barbadoes aloes. It is readily procured by making a decoction in water acidified with a little hydrochloric acid, after some hours pouring off the precipitated resin and evaporating the liquid to a syrup. The aloin crystallizes out in a day or two. Barbaloin yields, by the action of bromine and chlorine, substitution compounds. Nitric acid dropped upon it produces a red colour, which soon fades. Boiled for some time with strong nitric acid barbaloin gives, together with oxalic and picric acids, a yellow substance, chrysammic acid, which furnishes beautiful red salts (Tilden). Anthracene ($\text{C}_{14}\text{H}_{10}$) has been obtained by deoxidation of barbaloin.

Nataloin.—This body was discovered by Flückiger in Natal aloes. It crystallizes readily in rectangular plates, either from spirit or from water. No bromine or chlorine substitution derivatives have yet been formed, but an acetyl compound has been analysed (Tilden). Nataloin moistened with nitric acid gives a red coloration which does not fade. When boiled with nitric acid it yields no chrysammic, but only oxalic and picric acids.

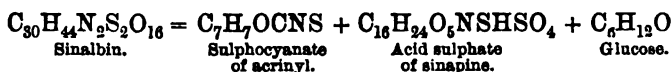
Socaloin or Zanaloin.—Histed and Flückiger have shown that Socotrine or Zanzibar aloes yields an aloin distinct from

those just described. It forms tufted acicular prisms. Nitric acid scarcely alters the colour of socaloin. Neither socaloin nor barbaloin afford any colour when vapour from a glass rod moistened with nitric acid is brought near to a drop of oil of vitriol containing a minute fragment of the aloin, while nataloin gives rise to a blue coloration.

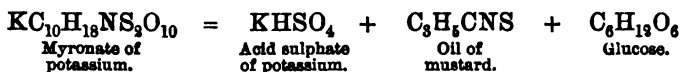
Analysis.—To Aloin or powdered aloes on a white plate add strong nitric acid. *No colour* = Socaloin. *Crimson colour* = Nataloin or Barbaloin. To another portion add strong sulphuric acid and vapour of nitric acid. *A blue colour* = Nataloin. *No blue colour* = Barbaloin.

Formulae.—The reactions of these bodies seem to indicate that they are complex phenols. Phenol being the phenyl hydrate, C_6H_5HO , and cresol being methyl-phenol, $C_6H_4CH_3HO$, the aloins may possibly have a similar constitution,—that is, they may be the hydrates of radicals in which part of the hydrogen is replaced by groups of atoms. Since Tilden made this suggestion, E. von Sommaruga and Egger (*"Pharmacographia"*) have arrived at the conclusion that the aloins form an homologous series, and that they have the composition indicated in the following formulae:—Socaloin, $C_{15}H_{16}O_7$; Nataloin, $C_{16}H_{18}O_7$; Barbaloin, $C_{17}H_{20}O_7$. Tilden's more recent experiments indicate, however, that barbaloin ($C_{16}H_{18}O_7$) and socaloin ($C_{16}H_{18}O_7$) are isomeric in the anhydrous state, but that socaloin and its derivatives in the hydrous condition contain more water of crystallization than barbaloin. Nataloin ($C_{16}H_{18}O_7$) seems to be isomeric with the others, but is less soluble, and does not yield either chrysammic acid or chloro- or bromo-derivatives ($C_{16}H_{15}Cl_3O_7$; $C_{16}H_{15}Br_3O_7$). The acetyl-derivatives appear to have the formula $C_{16}H_{15}(C_2H_3O)_3O_7$.

In the series C_nH_{2n-1} we have the univalent radical *allyl* (C_3H_5), whose sulphide [$(C_3H_5)_2S$] is *essential oil of garlic*, and sulphocyanate (C_3H_5CyS) the chief part of the *essential oil of mustard*, the body to which mustard owes its power of inducing inflammatory action on the skin (*"Mustard Poultice"* and *Charta Sinapis*, B. P.). *Mustard (Sinapis*, B. P.) is a powdered mixture of black or, rather, reddish-brown and white mustard-seeds. The white mustard-seed contains *sinalbin* ($C_{30}H_{44}N_2S_2O_{16}$), a glucoside which, in contact with the myrosin in an aqueous extract of mustard, yields the sulphocyanide of a peculiar radical *acrinyl*, a body which forms part of the essential oil of mustard paste.



The black contains the albumenoid ferment *myrosin*, resembling the *emulsin* of almonds (p. 491) and *myronate* of potassium, or *sinigrin*. The latter is the body which, under the influence of the former, yields the chief part of the pungent oil of mustard paste.



Crude oil of mustard often contains *cyanide of allyl*, $\text{C}_3\text{H}_5\text{CN}$.

In the Pharmacopœia of India the seed of *Sinapis juncea*, *Rai* or *Indian Mustard Plant*, is official in addition to that of *S. alba* and *S. nigra*. It is the common mustard of warm countries. It does not differ chemically from other mustard. Allyl compounds are also met with in several other cruciferous and liliaceous plants.

In the C_nH_{2n} series occurs *ethylene* or *olefiant gas* (C_2H_4), the chief illuminating constituent of coal-gas (made by heating spirit of wine with twice its volume of strong sulphuric acid), a bivalent radical, the alcohol of which is *glycol* ($\text{C}_2\text{H}_4\text{.2HO}$); and in the $\text{C}_n\text{H}_{2n-1}$ the trivalent hypothetical radical *glyceryl* (C_3H_5), the hydrate of which ($\text{C}_3\text{H}_5\text{.3HO}$) is *glycerine*. The homologues of glycol (*e. g.* propylene glycol and butylene glycol) are termed *glycols*, the homologues of glycerine *glycerines*. By oxidation of glycols, acids of the lactic series are formed (glycollic, lactic, and oxybutyric acids); just as by oxidation of ordinary alcohols acids of the acetic series are formed. It will be noticed that the chemical composition of the radicals glyceryl and allyl is identical (C_3H_5); but the former is trivalent and the latter univalent; hence they probably differ in physical constitution: they are isomeric, possibly polymeric, with each other. From a glyceryl compound (glycerine), however, an allyl salt (iodide) can be produced. By distilling a mixture of glycerine and diiodide of phosphorus, iodide of allyl ($\text{C}_3\text{H}_5\text{I}$) is obtained; and on digesting this with sulphocyanate of potassium, the sulphocyanate of allyl, or *artificial oil of mustard*, results, identical with the chief constituent of natural oil.

GLYCERINE.—Glycerine, or Glyceric Alcohol ($\text{C}_3\text{H}_5\text{.3HO}$), is the hydrate of the basylous radical of most oils and fats, bodies which are mainly oleates, palmitates, and stearates of

glyceryl (glycyl or propenyl). When these substances are heated with metallic hydrates (even with water—hydrate of hydrogen—at a temperature of 500° F. or 600° F.) double decomposition occurs, oleate, palmitate, or stearate of the metal is formed, and hydrate of glyceryl or glycerine. Hence glycerine is a by-product in the manufacture of soap, hard candles, and lead-plaster (*vide* Index).

Properties.—Glycerine is viscid when pure, specific gravity 1.28 (not below 1.25, B. P.), has a sweet taste, is soluble in water or alcohol in all proportions. It has remarkable powers as a solvent, is a valuable *antiseptic* even when diluted with 10 parts of water, and useful as an emollient. In vacuo it may be distilled unchanged, but under ordinary atmospheric pressure it is decomposed by heat. From damp air glycerine absorbs moisture slowly but in considerable proportions. Perfectly pure and anhydrous glycerine, at a few degrees below the freezing point of water, sometimes solidifies to a mass of crystals.

Tests.—Heat one or two drops of glycerine in a test-tube, alone or with strong sulphuric acid, acid sulphate of potassium, or other salt powerfully absorbent of water; vapours of acrolein (from *acer*, sharp, and *oleum*, oil) are evolved, recognised by their powerfully irritating effects on the eyes and respiratory passages. If the glycerine be in solution, it must be evaporated as low as possible before applying this test.

Add a few drops of the fluid suspected to contain the glycerine to a little powdered borax; stir well together; dip the looped end of a platinum wire into the mixture and expose to an air-gas flame; a deep green colour is produced (Senier and Lowe).

The glycerine liberates boracic acid, and it is the latter which colours the flame. Ammoniacal salts, which similarly affect borax, must first be got rid of by boiling with solution of carbonate of sodium. Liquids containing much indefinite organic matter must sometimes be evaporated to dryness, the residue extracted by alcohol, and the latter tested for the glycerine. To detect traces, liquids must be concentrated.

Besides glycerine itself (*Glycerinum*, B. P.), there are several official preparations of glycerine—solutions of carbolic, gallic and tannic acids and borax in glycerine, and a sort of

mucilage of starch in glycerine (*Glycerinum Acidi Carbolici*, *Glycerinum Acidi Gallici*, *Glycerinum Acidi Tannici*, *Glycerinum Boracis*, and *Glycerinum Amyli*).

QUESTIONS AND EXERCISES.

794. Give the formulæ of some compounds of radicals which have the general formulæ C_nH_{2n-7} , C_nH_{2n-1} , C_nH_{2n-1}' , and C_nH_{2n}'' ,—e.g., benzol, essential oil of mustard, glycerine, and glycol.

795. State the difference in composition of natural and artificial oil of bitter almonds.

796. How is the so-called artificial oil of bitter almonds prepared?

797. What are the uses, composition, source, and properties of Carbolic Acid?

798. State the characters by which carbolic acid is distinguished from Creasote.

799. Draw out an equation explanatory of the production of aniline.

800. Mention the chief properties of Glycerine.

801. What is the specific gravity of glycerine?

802. By what test is glycerine recognised?

803. Enumerate some official preparations in which glycerine is employed as a solvent.

ALBUMENOID SUBSTANCES.

Albumen.—Agitate, thoroughly, white of egg (*Albumen Ovi*, B. P.) with water, and strain or pour off the liquid from the flocculent membranous insoluble matter. One white to 4 oz. of water forms the "Solution of Albumen," B. P.

Test.—Heat a portion of this solution of albumen to the boiling-point; the albumen becomes insoluble, separating in clots or coagula of characteristic appearance.

Other Reactions.—Add to small quantities of aqueous solution of albumen solutions of corrosive sublimate, nitrate of silver, sulphate of copper, acetate of lead, alum, perchloride of

tin; the various salts not only coagulate but form insoluble compounds with albumen. Hence the value of an egg as a temporary antidote in cases of poisoning by many metallic salts, its administration retarding the absorption of the poison until the stomach-pump or other means can be applied. Sulphuric, nitric, and hydrochloric acids precipitate albumen; the coagulum is slowly redissolved by aid of heat, a brown, yellow, or purplish-red colour being produced. Neither acetic, tartaric, nor organic acids generally, except gallo-tannic, coagulate albumen. Alkalies prevent the precipitation of albumen.

Yolk or Yelk of Egg (Ovi Vitellus, B. P.) contains only 3 per cent. of albumen—the white $12\frac{1}{2}$. The yolk also contains 30 per cent. of yellow fat and 14 of casein.

Albumen is met with in large quantity in the serum of blood, in smaller quantity in chyle and lymph, and in the brain, kidneys, liver, muscles, and pancreas. It is not a normal constituent of saliva, gastric juice, bile, or mucus, but occurs in those secretions during inflammation. It is found in the urine and fæces only under certain diseased states of the system.

The cause of the coagulation of albumen by heat has not yet been discovered.

Albumen has never been obtained sufficiently pure to admit of its composition being expressed by a trustworthy formula; Gerhardt regarded it as a sodium compound ($\text{HNaC}_{72}\text{H}_{110}\text{N}_{18}\text{SO}_{22}, \text{H}_2\text{O}$).

Egg-albumen, and, to some extent, blood-albumen, is largely used by calico-printers as a vehicle for colours, serving also, when dry, as a glaze. Curriers prize egg-oil for softening leather.

Albumen coagulated by heat is said to be recoverable in a scarcely altered fluid condition by contact with a dilute aqueous solution of a very small proportion of pepsin.

Fibrin, Casein, Legumin.

Fibrin is the chief constituent of the muscular tissue of animals. It occurs in solution in the blood; and its spontaneous solidification or coagulation is the cause of the clotting of blood shortly after being drawn from the body—a phenomenon which cannot at present be explained satisfactorily.

Fibrin may be obtained by whipping fresh blood with a bundle of twigs, separating the adherent fibres, and washing in water till colourless. It may be dried or kept under spirit of wine.

Average Composition of Blood (in 1000 parts).

(Compiled by Kirkes.)

	Water	784
	Albumen	70
	Fibrin	2·2
	Red corpuscles (dried)	130
Fatty matters.	Cholesterin	0·08
	Cerebrin.	0·40
	Serolin	0·02
	Oleic and margaric acids	1·4
	Volatile and odorous fatty acid	
	Fat containing Phosphorus	
Inorganic salts.	Chloride of sodium	3·6
	Chloride of potassium	·35
	Phosphate of sodium (Na_3PO_4)	·2
	Carbonate of sodium	·82
	Sulphate of sodium	·23
	Phosphates of calcium and magnesium	·25
	Oxide and phosphate of iron	·50
	Extractive matters, biliary colouring-matter, gases, and accidental substances	6·4

1000·

Percentage proportion of the *chief* constituents of Blood.

Water	78·4
Red corpuscles (solid residue)	13·0
Albumen of serum	7·0
Inorganic salts	·603
Extractive, fatty, and other matters	·777
Fibrin	·22

100·

Casein occurs in Cow's Milk (*Lac*, B. P.) to the extent of 3 per cent., dissolved by a trace of alkaline salt. Its solution does not spontaneously coagulate like that of fibrin, nor by heat like albumen; but acids cause its precipitation from milk in the form of a curd (cheese) containing the fat- (butter-) globules previously suspended in the milk, a clear yellow

liquid (or whey) remaining. *Curd*s and *whey* are also produced on adding to milk a piece, or an infusion, of *rennet*, the salted and dried inner membrane of the fourth stomach of the calf. The exact action of rennet is not known. Respecting rennet Soxhlet says: "60 to 80 grammes of calf's stomach steeped for five days in 1 litre of a 5 per cent. solution of common salt at ordinary temperatures, yield a solution of which 1 vol. will coagulate 10,000 vols. of new milk at a temperature of 95° F. in forty minutes. If the filtered solution is treated with 60 to 90 grammes more of stomach a solution of double strength is obtained; another repetition gives a solution three times the strength of the original one. To prevent decomposition, about 0.3 per cent. of thymol may be added to the concentrated rennet extract solution. Possibly a slight taste due to this may be detected in the finest cheese, but for the same reason oil of cloves is much more objectionable. Boric acid is on all accounts the best antiseptic to employ, and solutions to which it has been added may be kept in covered vessels for months. All extract solutions lose strength on keeping; during the first two months the solution may become 30 per cent. weaker, then the strength remains nearly constant for eight months in the case of a solution of 1 in 18,000. Alcohol is almost as good an antiseptic as boric acid, if the solution be preserved in well-stoppered flasks."

Average Composition of 1000 parts of Milk.

	Specific gravity.	Water.	Solid constituents.	Casein and extractive.	Sugar.	Butter.	Salts.
Woman. . .	1.033	889	111	40	42	27	2
Cow. . . .	1.032	876	124	40	47	30	7

Specific gravity alone, as taken by the form of hydrometer termed a *lactometer*, or even by more delicate means, is of little value as an indication of the richness of milk, the butter and the other solids exerting an influence in opposite directions. Good cow's milk affords from 10 to 12 per cent. by volume of cream, and 3 to 3½ per cent. of butter. The water of milk seldom varies more than from 87 to 88 per cent., and the solid constituents from 13 to 12. Indeed, excluding its butter, milk is curiously regular in composition. The non-

fatty solids in the mixed milk of a herd or dairy of healthy cows is almost a constant quantity, namely 9·3 per cent. A lower proportion of non-fatty solids in a sample of milk points to the addition of water. Thus, supposing that 100 grains of a specimen of milk evaporated to dryness, and all butter extracted from the residue by ether, yielded a non-fatty residue of 7·44 grains, the specimen would probably be four-fifths milk and one-fifth water. For if (by rule of three) 9·3 indicate 100, then 7·44 indicate 80. Occasionally, under exceptional circumstances, a sample of genuine milk might be slightly poorer than that from a healthy herd, and therefore, for legal purposes, a standard of 9 per cent. by weight of non-fatty solids and 2·5 per cent. of butter-fat has been proposed. Only in the rare cases of milk containing an unusually large proportion of butter-fat would any milk yielding less than 9 per cent. of non-fatty solids be regarded as genuine. And, again, no milk would be considered genuine, under this standard, if it yielded less than 2·5 per cent. of fat, not even in the rare case of its containing an unusually large proportion of real non-fatty milk-solids. Half-starved cows might yield milk below these standards; but such milk could scarcely be considered to be genuine, or better fitted for food than milk watered *after* leaving the cow.

Under the microscope milk is seen to consist of minute corpuscles floating in a transparent medium. These corpuscles consist of the fatty matter (butter), said to be contained in a filmy albumenoid envelope. The fat is fluid at the normal temperature of the animal, and remains so until the milk is well agitated by churning or otherwise, or until the milk is frozen.

Legumin or *vegetable casein* is found in most leguminous seeds, such as sweet and bitter almonds. Peas contain about 25 per cent. of legumin.

Vegetable albumen is contained in many plant-juices, and is deposited in flocculi on heating such liquids. *Vegetable fibrin* is the name given by Liebig and Dumas to that portion of the gluten of wheat which is insoluble in alcohol and ether (*vide* p. 474).

Albumenoid substances are nearly identical in percentage composition. Albumen (and fibrin) contains 53·5 of carbon, 7 of hydrogen, 15·5 of nitrogen, 22 of oxygen, 1·6 of sulphur, and ·4 of phosphorus. Casein contains no phosphorus. These three bodies are often termed the *plastic elements of nutrition*,

under the assumption that animals directly assimilate them in forming muscles, nerves, and other tissues,—starch, sugar, and similar matter forming the *respiratory* materials of food, because more immediately concerned in keeping up the temperature of the body by the combustion going on between them, and their products, and the oxygen of the air in the blood.

The whole of the organic nitrogen in food must not however be regarded as representing true albumenoids, some existing as amidic and similar compounds, bodies having a simplicity of composition characteristic of the *products* of physiological action on food, rather than that complexity of composition characteristic of true nutrients. Albumenoids in decomposing yield much fatty as well as other substances. Possibly a portion, at least, of the *adipocere* (*adeps*, fat; *cera*, wax), or *corpse-fat*, characteristic of the remains of buried animals is thus derived.

Musk (*Moschus*, B. P.), “the inspissated and dried secretion from the preputial follicles of *Moschus moschiferus*” (the Musk-Deer), is a mixture of albumenoid, fatty, and other animal matters with a volatile odorous substance of unknown composition.

GELATIGENOUS SUBSTANCES.

These nitrogenous bodies differ, chemically, from the albumenoid in containing less carbon and sulphur and more nitrogen. They are contained in certain animal tissues, and on boiling with water yield a solution which has the remarkable property of solidifying to a jelly on cooling. The tendons, ligaments, bones, skin, and serous membranes afford *gelatine* proper; the cartilages give *chondrine*, which differs from gelatine in composition and in being precipitated by vegetable acids, alum, and the acetates of lead. The purest variety of gelatine is *isinglass* (B. P.), “the swimming-bladder or sound of various species of *Acipenser*, Linn., prepared and cut in fine shreds.” Small quantities are more easily disintegrated by a file than a knife. Fifty grains dissolved in 5 ounces of distilled water forms the official “Solution of Gelatine,” B. P. *Glue* is an impure variety of gelatine, made from the trimmings of hides; *size* is glue of inferior tenacity, prepared from the parings of parchment and thin skins. “Among the

varieties of gelatine derived from different tissues and from the same sources at different ages, much diversity exists as to the firmness and other characters of the solid formed on the cooling of the solutions. The differences between isinglass, size, and glue, in these respects, are familiarly known, and afford good examples of the varieties called weak and strong, or low and high, gelatines. The differences are sometimes ascribed to the quantities of water combined in each case with the pure or anhydrous gelatine, part of which water seems to be intimately united with the gelatine; for no artificial addition of water to glue would give it the character of size, nor would any abstraction of water from isinglass or size convert it into the hard dry substance of glue. But such a change is effected in the gradual process of nutrition of the tissues; for, as a general rule, the tissues of an old animal yield a much firmer or stronger jelly than the corresponding parts of a young animal of the same species." (Kirke's Physiology.)

Gelatine appears to unite chemically with a portion of the water in which it is soaked when used for culinary and manufacturing purposes, for a solution of glue in hot anhydrous glycerine does not yield an ordinary jelly on cooling. From its solution in water, gelatine is precipitated by alcohol, corrosive sublimate, perchloride of platinum, and by tannic acid. Its aqueous solution is not, like that of albumen, coagulated by heat. By prolonged ebullition its gelatinizing power is destroyed.

PEPSIN.

Pepsin (from *πέρω*, *pepto*, I digest) is a nitrogenous substance existing in the gastric juice, and as a viscid matter in the *peptic* glands and on the walls of the stomachs of animals. To isolate it, the mucous membrane of the stomach (of the hog, sheep, or calf, killed fasting) is scraped, and macerated in cold water for twelve hours; the pepsin in the strained liquid is then precipitated by acetate of lead, the deposit washed once or twice by decantation, sulphuretted hydrogen passed through the mixture of the deposit with a little water to remove the whole of the lead, and the filtered liquid evaporated to dryness at a temperature not exceeding 105° F. Pepsin is a powerful promoter of digestion; its solution is hence frequently termed *artificial gastric juice*. As met with in pharmacy its strength varies greatly. It is often prepared by simply mixing with starch the thick liquid obtained on

macerating the scraped stomach with water, and evaporating to dryness. (*Vide Pharmaceutical Journal*, 1865-66, p. 112, and 1871-72, pp. 785 and 843.) The official process (*Pepsin*, B. P.) simply consists in scraping the viscid pulp from the slightly washed inner surface of the stomach, and quickly evaporating it to dryness on glass or glazed earthenware at a temperature not exceeding 100° F. The product is powdered. "A light yellowish brown powder, having a faint, but not disagreeable odour, and a slightly saline taste, without any indication of putrescence. Very little soluble in water or spirit. Two grains of it with an ounce of distilled water, to which five minims of hydrochloric acid have been added, form a mixture in which 100 grains of hard-boiled white of egg, in thin shavings, will dissolve on their being digested together for about four hours at a temperature of 98°." The solvent or digestive action of pepsin on the albumenoids, etc., in the stomach results in a fluid termed *peptone*. It is thus that such food is prepared for conversion into blood. Artificial peptone is made by digesting blood fibrin with pepsin in very weak hydrochloric acid. Peptone is not readily coagulated by heat, and it freely diffuses through membranes. It appears to be isomeric with albumen. Some vegetables, notably the leaves of the papaw tree, *Carica papaya*, appear to contain a principle, "papaine," analogous in properties to pepsin. According to Wurtz papaine is an albumenoid.

(For a *résumé* of the different modes of preparing pepsin, see an article by Petit in the *Pharmaceutical Journal* for July 17, 1880.)

PANCREATIN.

The pancreas (or "sweetbread") secretes a colourless fluid which contains $1\frac{1}{2}$ to $2\frac{1}{2}$ per cent. of an albumenoid substance, which has the power of converting starch into sugar and, especially, of emulsifying fat. It may be precipitated by chloride of sodium from an acidulated infusion of the pancreas. It is soluble in cold water. An extremely small proportion emulsifies a large volume of fat.

QUESTIONS AND EXERCISES.

804. In what form is Albumen familiar?

805. Name the chief test for albumen.

806. Why is the administration of albumen useful in cases of poisoning?

807. Mention the points of difference between yolk and white of egg.

808. From what sources other than egg may albumen be obtained?

809. In what respects does fibrin differ from albumen?

810. Enumerate the chief constituents of blood.

811. How may fibrin be obtained from blood?

812. State the differences between casein, fibrin, and albumen.

813. What are the relations of cream, butter, curds and whey, and cheese, to milk?

814. Describe the microscopic appearances of blood and of milk.

815. How much cream should be obtained from good milk?

816. What is the percentage of water in genuine milk?

817. Name sources of vegetable casein and vegetable albumen.

818. Give the percentage of nitrogen in albumenoid substances.

819. Describe the chemical nature of Musk.

820. In what lie the peculiarities of Gelatine?

821. To what extent do isinglass, glue, and size differ?

822. Whence is Pepsin obtained?

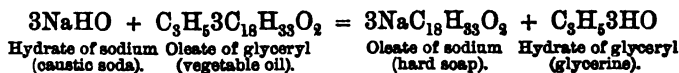
823. How is pepsin prepared?

FATTY BODIES.

SOAPS, SOLID FATS, FIXED OILS, VOLATILE OILS, CAMPHORS.

General relations.—Oils and fats are, apparently, almost as simple in constitution as ordinary inorganic salts. Just as acetate of potassium ($\text{KC}_2\text{H}_3\text{O}_2$) is regarded as a compound of potassium (K) with the characteristic elements of all acetates ($\text{C}_2\text{H}_3\text{O}_2$), so soft soap is considered to be a compound of potassium (K) with the elements characteristic of all oleates ($\text{C}_{18}\text{H}_{33}\text{O}_2$), and hence is chemically termed oleate of potassium ($\text{KC}_{18}\text{H}_{33}\text{O}_2$). Olive-oil, from which soap is commonly prepared, is mainly oleate of the trivalent radical *glyceryl* (C_3H_5), the formula of pure fluid oil being $\text{C}_3\text{H}_5\text{C}_{18}\text{H}_{33}\text{O}_2$, and its name *oleine*. The formation of a soap therefore, on bringing together oil and a moist oxide or hydrate, is a simple case of double decomposition (or, rather, metathesis), as seen already

in connection with lead plaster (p. 242), or in the following equation relating to the formation of common hard soap :—



Berthelot has succeeded in preparing oil artificially from oleic acid and glycerine; and it is said to be identical with the pure oleine of olive- and of other fixed oils. Hard fats chiefly consist of *stearine*—that is, of tristearate of glyceryl ($\text{C}_3\text{H}_5\text{3C}_{18}\text{H}_{35}\text{O}_2$). Mr. Wilson, of Price's Candle Company, obtains stearic and oleic acids and glycerine by simply passing steam, heated to 500° or 600° F., through melted fat. Both the glycerine and fat-acids distil over in the current of steam, the glycerine dissolving in the condensed water, the fat-acids floating on the aqueous liquid.

The author finds that oleic acid readily combines with alkalis and most of the metallic oxides or hydrates, forming oleates which are soluble in fats. In this way active medicines may be administered internally in conjunction with oils, or externally in the form of ointments.

As regards the conversion of oily substances into emulsions resembling the common natural emulsion, milk, Gregory states that three drachms of gum acacia in fine powder are necessary to emulsify one ounce of any of the volatile oils, and that a little less (about two drachms) will answer for the fixed oils and balsams. To this quantity of gum four drachms and a half of water must be added (no more and no less). Either the water or the oil may be added first to the gum, but it is quickest to add the oil first, and well triturate before adding the water.

SOAPS.—Olive-oil boiled with solution of potash yields potassium soap, or *soft soap* (*Sapo Mollis*, B. P.); with soda, sodium soap, or *hard soap* (*Sapo Durus*, B. P.); mixed with ammonia, an ammonium soap (*Linimentum Ammoniae*, B. P.); and with lime-water, calcium soap (*Linimentum Calois*, B. P.),—all oleates, chiefly, of the respective basylous radicals. Their mode of formation is indicated in the foregoing equation. The alkali soaps are soluble in alcohol, the others insoluble. A green soap, much used on the continent of Europe, and indeed, official in Germany (*Sapo Viridis*), is made by adding

indigo to ordinary soft soap; the yellow colour of the soap yielding with the indigo a greenish compound. The official characters of Hard Soap are:—"greyish-white, dry, inodorous; horny and pulverizable when kept in dry warm air; easily moulded when heated; soluble in rectified spirit; not imparting an oily stain to paper: incinerated it yields an ash which does not deliquesce." And of Soft Soap:—"yellowish-green, inodorous, of a gelatinous consistence; soluble in rectified spirit; not imparting an oily stain to paper: incinerated it yields an ash which is very deliquescent." Curd Soap (*Sapo Animalis*, B. P.) is "a soap made with soda and a purified animal fat, consisting principally of stearin." It will, of course, chiefly contain stearate of sodium. In pharmacy it is often advantageously employed instead of the "hard soap."

The hard soap met with in trade is made from all varieties of oil, the commoner kinds being simply the product of the evaporated mixture of oil and alkali; while the better sorts have been separated from alkaline impurities and the glycerine by the addition of common salt to the liquors, which causes the precipitation of the pure soap as a curd. Potash soap is not so readily precipitable by salt.

Bile, the gall of the ox (*Bos taurus*, Linn.), freed from mucus by agitating with twice its bulk of rectified spirit (in which mucus is insoluble), filtering, and evaporating, yields the official Purified Ox-Bile (*Fel Bovinum Purificatum*, B. P): the latter has a resinous appearance, but is chiefly composed of two crystalline substances having the constitution of a soap; the one is termed *taurocholate of sodium* ($\text{NaC}_{26}\text{H}_{44}\text{NO}_7$ S), the other is *glycocholate*, or simply *cholate*, of sodium ($\text{NaC}_{26}\text{H}_{43}\text{NO}_6$). Both taurocholates and glycocholates are conjugated bodies readily yielding, the former *cholic* or *cholalic acid* ($\text{HC}_{24}\text{H}_{39}\text{O}_5$) and *taurine* ($\text{C}_9\text{H}_7\text{NO}_3\text{S}$), the latter *cholalic* and *glycocine* or *glycocoll* ($\text{C}_9\text{H}_5\text{NO}_3$), a soluble crystalline body having interesting physiological relations, inasmuch as it is obtainable from gelatine (hence the name glycocoll or sugar of gelatine, from γλυκὺς, *glucūs*, sweet, and κόλλα, *kolla*, glue) and from hippuric acid. The presence of bile in a liquid, such as urine, is detected by Pettenkofer's test. The fluid is gradually mixed with half its bulk of strong sulphuric acid in a test-tube, rise of temperature being prevented by partial immersion

of the tube in water. A small quantity of powdered white sugar is then introduced and well mixed with the acid liquid, and more sulphuric acid then poured in; as the temperature rises a reddish or violet coloration is produced. The cholic acid liberated in the reaction furnishes the colour.

SOLID FATS.—1. *Lard* (*Adeps Preparatus*, B. P.) is the purified internal fat of the abdomen of the hog—the perfectly fresh *omentum* or *flare*, washed, melted, strained, and dried. 2. *Benzoated Lard* (*Adeps Benzoatus*, B. P.) is prepared lard heated over a water-bath with benzoin (ten grains per ounce), which communicates an agreeable odour and prevents or retards rancidity. Purified lard is a mixture of oleine and stearine; *margarine*, the margarate of glyceryl, was formerly supposed to be a constituent of lard and other soft fats, but is now regarded as a mere mixture of *palmitine* (the chief fat of palm-oil) and stearine. 3. *Yellow Wax* (*Cera Flava*, B. P.), the prepared honeycomb of the Hive-Bee, and the same bleached by exposure to sunlight. 4. *White Wax* (*Cera Alba*, B. P.), according to Brodie, is chiefly a mixture of *Cerotic Acid* ($\text{HC}_{27}\text{H}_{53}\text{O}_2$), *Palmitate of Melissyl* ($\text{C}_{30}\text{H}_{61}\text{C}_{16}\text{H}_{31}\text{O}_2$), and about 5 per cent. of *Ceroleine*, the body to which the colour, odour, and tenacity of wax are due. *Ceresine*, the purified native *ozokerite* of Galicia, is a solid hydrocarbon largely used as a substitute for beeswax, especially in Russia. It differs from beeswax chemically in being almost unattacked by warm oil of vitriol. 5. *Spermaceti* (*Cetaceum*, B. P.) is the *palmitate of cetyl* ($\text{C}_{16}\text{H}_{33}\text{C}_{16}\text{H}_{31}\text{O}_2$) or *cetine*; when saponified it yields not glycerine, the hydrate of glyceryl ($\text{C}_3\text{H}_5\text{O}$), but *ethyl*, the hydrate of cetyl ($\text{C}_{16}\text{H}_{33}\text{HO}$); it is the solid crystalline fat accompanying *sperm-oil* in the head of the spermaceti-whale. 6. *Suet*, the internal fat of the abdomen of the sheep, purified by melting and straining, forms the official *Prepared Suet* (*Sevum Preparatum*, B. P.); it is almost exclusively composed of stearine ($\text{C}_3\text{H}_7\text{C}_{18}\text{H}_{35}\text{O}_2$). 7. Expressed oil of *nutmeg* (*Oleum Myristicæ Expressum*, B. P.), commonly but erroneously termed *Oil of Mace*, is a mixture of a little volatile oil with much yellow and white fat, the latter is *myristin* or *myristate* of glyceryl ($\text{C}_3\text{H}_5\text{C}_{14}\text{H}_{27}\text{O}_2$). 8. Oil of *theobroma*, or *Cacao-butter* (*Oleum Theobromæ*, B. P.), chiefly theobromate of glyceryl (Kingzett), is a solid product of the roasted and roughly crushed seeds or *cocoa nibs* of the *Theobroma cacao*. They contain from one-fourth to one-half of this fat. (These also furnish, when ground, *flake cocoa*;

or when ground and much sweetened, *chocolate*; or, with farina and some sugar, *cocoa*; or with a portion of the butter extracted, "*cocoatina*," etc.) 9. *Cocoa-nut* oil or butter, a soft fat largely contained in the edible portion of the nut of *Cocos nucifera*, or common cocoa-nut of the shops, a body containing glyceryl united with no less than six different univalent acidulous radicals, namely, the caproic ($C_6H_{11}O_2$), caprylic ($C_8H_{15}O_2$), rutic ($C_{10}H_{19}O_2$), lauric ($C_{12}H_{23}O_2$), myristic ($C_{14}H_{27}O_2$), and palmitic ($C_{16}H_{31}O_2$)—radicals which, like some from common resin, when united with sodium, form a soap differing from ordinary hard soap (oleate of sodium) by being tolerably soluble in a solution of chloride of sodium; hence the use of cocoa-nut oil and resin in making *marine soap*, a soap which, for the reason just indicated, readily yields a lather in sea-water. 10. *Kokum Butter*, *Garcinia Oil*, or Concrete Oil of Mangosteen, a whitish or yellowish-white fat obtained from the seeds of *Garcinia indica* or *G. purpurea*, is composed of stearine, myristicine, and oleine. It is recognised officially in the Pharmacoposia of India (*Garcinia purpureae Oleum*).

Butter yields $87\frac{1}{2}$ per cent. of fat acids by saponification and decomposition of the soap by acid. Other animal fats, with which butter is likely to be adulterated, yield about $95\frac{1}{2}$. Hence the percentage of fat acid yielded by a suspected sample of butter indicates not only purity or the opposite but extent of sophistication.

FIXED OILS.—*Fixed* and *Volatile* oils are naturally distinguished by their behaviour when heated; they also generally differ in chemical constitution—a fixed oil being, apparently, a combination of a basylous with an acidulous radical, while a volatile oil is more commonly a neutral hydrocarbon or the same oxidized.

Drying and Non-drying Oils.—Among fixed oils, most of which are oleate with a little palmitate and stearate of glyceryl, a few, such as, 1, *linseed-oil* (*Oleum Lini*, B. P. contained in *Lini Semina*, B. P., the ground residue of which, after removal of much oil, is *linseed meal*, *Lini Farini*, B. P.), and, 2, *cod-liver oil* (*Oleum Morrhue*, B. P.), and, to some extent, *castor* and *croton*, are known as *drying oils*, from the readiness with which they absorb oxygen and become hardened to a resin. Linseed commonly contains 37 or 38 per cent. of oil; 25 to 27 per cent. is obtained by submitting the ground seeds to hydraulic pressure, 10 or 12 per cent. re-

maining in the residual oil-cake. Boiled oil is linseed-oil which has been boiled with oxide of lead. This treatment increases the already great tendency of linseed-oil to resinify, forming *linozyn* ($C_{33}H_{54}O_{11}$) on exposure to air. The drying oils appear to contain linoleine, an oily body distinct from oleine. Among the *non-drying oils* are the following:—

3. *Almond-oil*, indifferently yielded by the bitter (*Amygdala Amara*, B. P.) or sweet seed (*Amygdala Dulcis*, B. P.) to the extent of 45 and 50 per cent. respectively.
4. *Croton-oil* (*Oleum Crotonis*, B. P.). Geuther states that no such acid as crotonic is obtainable from croton-oil, but acetic, butyric, valerianic, and higher members of the oleic series, together with *tiglic acid*, $HC_5H_7O_2$.
5. *Lycopodium*, a yellow powder composed of the spores of the common Club-Moss (*Lycopodium clavatum*), contains a large proportion of a very fluid fixed oil.
6. *Olive-oil* (*Oleum Olivæ*, B. P.), already noticed.
7. *Castor-oil* (*Oleum Ricini*, B. P.), chiefly a *ricinoleate of glyceryl* ($C_3H_5 \cdot 3C_{18}H_{33}O_2$) or ricinoleine, a slightly oxidized oleine, soluble, unlike most fixed oils, in alcohol and in glacial acetic acid. Castor-oil seeds are stated, by Tuson, to contain an alkaloid, *ricinine*. It possesses no purgative property.
8. Oil of *male fern* (*Filix Mas*, B. P.), a vermifuge obtained by exhausting the rhizome with ether and removing the ether by evaporation—a dark-coloured oil containing a little volatile oil and resin, and officially termed an extract (*Extractum Filicis Liquidum*, B. P.). Its chief active constituent appears to be *filicic acid*, $C_{14}H_{18}O_5$.
9. Fixed oil of *mustard*, a bland inodorous, yellow or amber oil, yielding by saponification and action of sulphuric acid, glycerine, oleic acid, and *erucic acid* ($HC_{22}H_{41}O_2$) (Darby).
10. *Arachis-oil* (*Oleum Arachis*, P. I.) is found to the extent of 40 or 50 per cent. in the seeds of the *Arachis hypogæa* (P. I.), the Ground-nut or Earth-nut (so-called because the pod of the herb in the growth of its stalk downwards is forced beneath the surface of the ground and there ripens). It is chiefly olein, but contains hypogæin, palmitin, and arachin. The oil is largely used in India in the place of olive-oil, and is becoming much employed in Europe, especially for soap-making.
11. *Sesamé-oil* (Gingelly, Teal, or Benne Oil), from the seeds of *Sesamum indicum*, is also largely used in Europe. It has most of the characters of the best olive-oil.
12. *Shark-liver oil*, from *Squalus carcharias* (*Oleum Squalæ*, P. I.), is used to some extent as a substitute for cod-liver oil in India.

Acids.	Formulae. Molec. Vol.=2.	Melting-point.		Boiling-point.		Whence obtained.
		°F.	°C*	°F	°C	
Formic . .	HC H O_2	47.3	8.5	212	100	{ Red ants; distillation of oxalic acid; and oxidation of amylaceous and other organic bodies.
Acetic . .	$\text{HC}_2\text{H}_3\text{O}_2$	62	16.7	244	118	{ Distillation of wood; oxidation of alcohol, etc.
Propionic .	$\text{HC}_3\text{H}_5\text{O}_2$	-58	-21	285	140.7	{ Fermentation of glycerin, etc.
Butyric . .	$\text{HC}_4\text{H}_7\text{O}_2$	32°	0	324	162.4	{ Butter; fermentation of lactic acid, etc.
Valerianic .	$\text{HC}_5\text{H}_9\text{O}_2$	3.2	-16	365	185	{ Valerian-root; oxidation of fousel oil.
Caproic . .	$\text{HC}_6\text{H}_{11}\text{O}_2$	28.4	-2	401	205	{ Butter.
(Enanthylic	$\text{HC}_7\text{H}_{13}\text{O}_2$	19	-10.5	435	224	{ Castor-oil by distillation, etc.
Caprylic .	$\text{HC}_8\text{H}_{15}\text{O}_2$	61.7	16.5	457	236	{ Butter; cocoa-nut oil.
Pelargonic .	$\text{HC}_9\text{H}_{17}\text{O}_2$	54.5	12.5	129	254	{ Leaves of the geranium.
Rutic. . .	$\text{HC}_{10}\text{H}_{19}\text{O}_2$	86	30	516	269	{ Butter; oil of rue by oxidation.
Lauric . .	$\text{HC}_{12}\text{H}_{23}\text{O}_2$	110	43	...	...	{ Cocoa-nut oil; berries of the bay tree.
Myristic .	$\text{HC}_{14}\text{H}_{27}\text{O}_2$	129	54	...	...	{ Nutmeg-butter; cocoa-nut oil, etc.
Palmitic .	$\text{HC}_{16}\text{H}_{31}\text{O}_2$	143.6	62	...	...	{ Palm-oil; butter; beeswax, etc.
Stearic . .	$\text{HC}_{18}\text{H}_{35}\text{O}_2$	44.4	69	...	...	{ Most solid animal fats.
Arachidic .	$\text{HC}_{20}\text{H}_{39}\text{O}_2$	167	75	...	...	{ Butter; oil of ground-nut.
Cerotic . .	$\text{HC}_{27}\text{H}_{55}\text{O}_2$	174	79	...	...	{ Beeswax.
Melissic . .	$\text{HC}_{20}\text{H}_{39}\text{O}_2$	195.8	91	...	...	{ Beeswax.

Their physical qualities and the formulæ of their acidulous radicals show that the fatty bodies are closely related, and indicate that the natural processes by which they are formed are probably as closely related. The above Table shows the homology of the fat-acids, and gives their names, formulæ, melting-points, boiling-points, and natural and artificial

* For the relation of degrees F. to C. *vide* Index, "Thermometers."

sources. (For further information respecting these acids and their isomers, the reader is referred to the Index, to works on "Organic Chemistry," to larger chemical works, or to Watts's "Dictionary of Chemistry.")

QUESTIONS AND EXERCISES.

824. Give a sketch of the general chemistry of fixed oils, fats, and soaps.

825. What is the difference between Hard and Soft Soap?

826. Which soaps are official?

827. Name the source of lard, and state how "Prepared Lard" is obtained.

828. State the composition of Beeswax.

829. In what does Spermaceti differ from other solid fats?

830. Mention the chief constituent of Suet.

831. Whence is Cacao-Butter obtained?

832. Why is *marine soap* so called? and from what fatty matter is it almost exclusively prepared?

833. What do you understand by *drying* and *non-drying* oils?

834. In what respect does Castor-Oil differ from other oils?

835. How is oil of male fern (*Ex. Filicis Liquidum*) prepared?

836. Mention the sources and formulæ of the following fatty acids:—formic, acetic, propionic, butyric, valerianic, caproic, cœnanthyllic, caprylic, pelargonic, and rutic.

VOLATILE OILS.—The *Volatile* or *Essential Oils* exist in various parts of plants, at first, probably, as mere combinations of carbon and hydrogen; but such *hydrocarbons* are prone to change when in contact with oxygen or moisture; hence these liquids, even when freshly obtained from the plants, and more especially as they occur in pharmacy, are usually mixtures of the liquid hydrocarbons or *elæoptens* (from *ἔλαιον*, *elæion*, oil, and *ὑπομαί*, *optomai*, I see) (pp. 552–557) with oxidized hydrocarbons, which are commonly solid or camphor-like bodies termed *stearoptens* (from *στέαρ*, *stear*, suet) (p. 558). The oils are also often associated with further oxidized bodies termed *resins* (p. 560). Of the hydrocarbons, those most commonly occurring are identical with or are isomers of that from oil of turpentine, and are known as *terpenes*. Tilden

separates these terpenes into two groups: a *turpentine group* of true terpenes ($C_{10}H_{16}$) whose members boil at 156° to 160° C., yield a nitroso-derivative melting at 129° , and form a solid crystalline hydrous terpene $C_{10}H_{20}O_3$, H_2O ; and an *orange group* ($C_{15}H_{24}$), polymerides of the true terpenes, whose members boil at 174° to 176° , yield a nitroso-derivative melting at 71° C., and form no solid hydrous compound. Kingzett's researches had previously pointed to a similar grouping, for he showed that turpentine and the other similar hydrocarbons ($C_{10}H_{16}$), and *cymene* ($C_{10}H_{14}$)—the probable nucleus of the terpenes—when mixed with water and subjected to a current of air, yielded to the water peroxide of hydrogen (not ozone), camphor and camphoric acid (the peroxide of hydrogen and camphoric acid possibly resulting from the action of water on a camphoric peroxide, $C_{10}H_{14}O_4$), while the second class ($C_{15}H_{24}$) yield no peroxide of hydrogen. The members of the $C_{10}H_{16}$ group are easily convertible into their polymers $C_{15}H_{24}$ and $C_{20}H_{32}$ by the action of heat, strong acids, etc. As regards either class, the members often differ from each other in their action on polarized light. Among the oxidized hydrocarbons occur several *aromatic aldehyds*. These, like ordinary aldehyds, form stable crystalline compounds with acid sulphite of sodium. On cooling a volatile oil, a stearopten often crystallizes out; or on distilling an oil, it remains in the retort, being less volatile than an elæopten. Volatile oils should, obviously, be preserved in well-closed bottles. Oxidation also proceeds more slowly in a cold than in a warm temperature.

(For a paper by Symes on the varying power of a large number of essential oils to twist a ray of plane polarized light, see the *Pharmaceutical Journal* for September 13, 1879.)

The process by which volatile oils are usually obtained from herbs, flowers, fruits, or seeds may be imitated on the small scale by placing the material (bruised cloves or caraways for instance) in a tubulated retort, adapting the retort to a Liebig's condenser, and passing steam, from a Florence flask, through a glass tube to the bottom of the warmed retort. The steam in its passage through the substance will carry the particles of oil over the neck of the retort into the condenser, and thence, liquefied and cooled, into the receiving vessel, where the oil will be found floating on the water. It may be col-

lected by running off the distillate through a glass funnel having a stopcock in the neck, or by letting the water from the condenser drop into an old test-tube which has a small hole in the bottom, or any similar tube placed in a larger vessel, the water and oil being subsequently run off separately from the tube as from a pipette. The water will in most cases be the ordinary official medicated water of the material operated on (*Aqua Aurantii Floris*, *Anethi*, *Oarui*, *Cinnamomi*, *Fœniculi*, *Menthæ Piperitæ*, *Menthæ Viridis*, *Pimentæ*, *Rosæ*—from *Rosæ Centifoliæ Petala*, B. P.,—*Sambuci*). Volatile oils, like fixed oils, stain paper; but the stain of the former is not permanent like that of the latter. Oils of lemon and orange are sometimes obtained by mere pressure of the rind of the fruit.

The presence of alcohol in an essential oil may be detected and its quantity estimated by slaking with an equal bulk of pure glycerine. The latter dissolves the alcohol and is augmented in volume according to the amount of alcohol present (Boettger). (For tests for the alcohol, see p. 506.)

A large number of volatile oils are employed in medicine, either in the pure state, in the form of saturated aqueous solution (medicated waters), solution in spirit of wine, 1 in 5 (*Essentia Anisi* and *Essentia Menthæ Piperitæ*, B. P.) and 1 in 50 (*Spiritus Cajuputi*, *Juniperi*, *Lavandulæ*, *Menthæ Piperitæ*, *Myristicæ*, *Rosmarini*), or as leading constituents in various barks, roots, leaves, etc. *Perfumes* ("scents" or "essences," including "Lavender-Water" and "Eau de Cologne") are for the most part solutions of essential oils in spirit of wine, or spirituous infusions of materials containing essential oils. The following oils are, directly or indirectly, official in the Pharmacopœias:—1. Volatile oil of *Bitter Almond* (p. 491). 2. Oil of the fruits of *Ajwain* or *Omum*, *Carum Ajowan*, or *Ptychotis Ajowan* (*Fructus Ptychotis*, P. I.), contains *cymol* or *cymene* ($C_{10}H_{14}$) and a *stearopten* (*Ajwain-ka-phul*, flowers of ajwain) identical with *thymol*, $C_{10}H_{14}O$. 3. Oil of *Dill* (*Oleum Anethi*, B. P.), a pale, yellow, pungent, acid liquid distilled from dill-fruit; it contains a hydrocarbon, *anethene* ($C_{10}H_{16}$), and an oxidized oil ($C_{10}H_{14}O$) identical with the carvol of oil of caraway (Gladstone). 4. Oil of *Aniseed* (*Oleum Anisi*, B. P.), a colourless or pale yellow liquid, of sweetish warm flavour, distilled in Europe

from the Anise-fruit (*Pimpinella anisum*), and in China from the fruit of Star-Anise (*Illicium anisatum*); it is a mixture of a hydrocarbon isomeric with oil of turpentine and *anethol*, a stearopten ($C_{10}H_{12}$) which crystallizes out at low temperatures. 5. Oil of *Chamomile* (*Oleum Anthemidis*, B. P.), a bluish or, when old, yellow oil, of characteristic odour and taste, distilled from chamomile-flowers (*Anthemidis flores*, B. P.). The official variety (*Anthemis nobilis*) yields about 0.2 per cent. of an oil composed of a hydrocarbon ($C_{10}H_{16}$) and an oxidized portion ($C_{10}H_{16}O_2$) which, heated with potash, gives *angelate of potassium* ($KC_8H_7O_2$), whence is obtained *angelic acid* ($HC_8H_7O_2$). According to Demarçay, Kopp, and Köbig, the oil is a mixture of the angelates of butyl and amyl and similar bodies. The flowers of another variety (*Matricaria chamomilla*) contain a stearopten ($C_{10}H_{16}O$) having the composition of laurel-camphor. 6. Oil of *Horse-radish-root* (*Armoracæ Radix*, B. P.) is, according to Hofmann, the sulphocyanate of butyl or tetryl (C_4H_9CNS): it is the chief active ingredient of *Spiritus Armoracæ Compositus*, B. P. 7. Oil of Bitter-Orange rind (*Aurantii Cortex*, B. P., *Aurantii Fructus*, B. P.), the flavouring constituent of the official syrup of the peel (*Syrupus Aurantii*, B. P.), and the oils of various species of *Citrus*, namely, 8, *lemon* (*Oleum Limonis*, B. P.), 9, *lime*, 10, *bergamot*, 11, *citron* and a variety of citron termed *cedra*, resemble each other in composition, all containing *hesperidene*, a hydrocarbon ($C_{15}H_{24}$), and a small quantity of oxidized hydrocarbons [$C_{10}H_{10}O_5$, $C_{15}H_{10}O$, and (Wright and Piesse) $C_{30}H_{30}O_5$]. Tilden states that lemon oil distilled from the fresh peel consists chiefly of a terpene, $C_{10}H_{16}$, boiling at $176^\circ C.$, with small quantities of a terpene boiling below 160° , and a hydrous terpene; the odour of the oil being due to the mixture. Expressed lime essence contains also a soft resin. 12. Oil of *Neroli* or *Orange-Flower*, the aqueous solution of which is official in the forms of water (*Aqua Aurantii Floris*, B. P.) and syrup (*Syrupus Aurantii Floris*, B. P.), contains a fragrant hydrocarbon ($C_{10}H_{16}$), colourless when fresh, but becoming red on exposure to light, and an inodorous oxidized hydrocarbon. Strong acids, especially nitric, attack the oil in Orange-flower water, colouring the fluid of a rose tint. 13. Oil of *Petit Grain*, distilled from the leaves and shoots of the Orange-tree, consists chiefly of a hydrocarbon apparently identical with that of oil of neroli. 14. The leaves of *Boldo* (*Peumus boldus*), a Chilean

shrub (tonic and hepatic), yield two per cent. of essential oil (and, according to Bourgon and Verne, an alkaloid, *boldine*). 15. Oil of *Buchu-leaves* (*Buchu Folia*, B. P.) consists of a hydrocarbon holding in solution a crystalline stearopten. Wayne has "more than once obtained from buchu a substance having the characters of salicylic acid." 16. Oil of *Cannabis indica*, see page 561. 17. Oil of (the lesser) *Cardamoms*, from the seeds of the capsules (*Cardamomum*, B. P.), is chiefly a hydrocarbon ($C_{10}H_{16}$) isomeric with oil of turpentine and a camphor resembling turpentine-camphor ($C_{10}H_{16}3H_2O$). 18. Oil of *Cajuput* (*Oleum Cajuputi*, B. P.) is a mobile bluish liquid, consisting chiefly of *hydrous cajuputene* or *cajuputol* ($C_{10}H_{16}, H_2O$). The latter, repeatedly distilled from phosphoric anhydride, yields cajuputene itself ($C_{10}H_{16}$), which has the odour of hyacinths. Fresh cajuput-oil has a green hue, which is perhaps transient, for the colour of the oil of trade is due to copper (Guibourt and Histed): certainly the green colouring-matter of pure cajuput-oil is organic. 19. Oil of *Caraway-fruit* (*Oleum Carui*, B. P.) is a mixture of *carvene* ($C_{15}H_{24}$) and *carvol* ($C_{10}H_{14}O$). 20. Oil of *Cloves* (*Oleum Caryophylli*, B. P.) and of *Pimento* (*Oleum Pimentæ*, B. P.), both heavier than water, contain a liquid hydrocarbon ($C_{20}H_{32}$), *eugenic acid* ($C_{10}H_{12}O_2$), a solid body, *eugenin*, isomeric with the eugenic acid, a second crystalline substance, *caryophyllin* ($C_{10}H_{16}O$), isomeric with common camphor, and a salicylic compound. 21. Oil of *Cascarilla-bark* (*Cascarillæ Cortex*, B. P.) has not been fully examined. 22. Oil of *Cinnamon-bark* (*Cinnamomi Cortex*, B. P.) and of *Cassia-bark* is mostly hydride of cinnamyl or cinnamic aldehyd (C_9H_7OH). Boiled with nitric acid it furnishes hydride of benzoyl (C_7H_5OH) and benzoic acid ($HC_7H_5O_2$), with chloride of lime yields benzoate of calcium ($Ca_2C_7H_5O_2$), and with caustic potash gives cinnamate of potassium ($KC_9H_7O_2$). The specific gravity of oil of cinnamon (*Oleum Cinnamomi*, B. P.) varies from 1.025 to 1.050. 23. Oil of *Citronella*, a *Grass Oil*, from *Andropogon nardus*, is chiefly composed of *citronellol* ($C_{10}H_{18}O$ and $C_{10}H_{18}O$, Wright), probably isomeric with the *absinthol* of wormwood (Gladstone). 24. Oil of *Copaiva* (*Oleum Copaibæ*, B. P.) and, 25, of *Cubebs* (*Oleum Cubebæ*, B. P.) are hydrocarbons having the formula $C_{15}H_{24}$. This *cubebene* is sometimes associated with a camphor, *hydrous cubebene* ($C_{15}H_{24}, H_2O$). Oil of cubebs also contains a small quantity of a terpene ($C_{10}H_{16}$). 26. Oil of *Coriander* (*Oleum Coriandri*, B. P.) seems

to have the composition of hydrous oil of turpentine ($C_{10}H_{16}H_2O$). 27. The fruits of *Oumin* or *Oummin* (*Cuminum cyminum*), an ingredient of many curry-powders, contains about 3 per cent., and those of *Water Hemlock* or *Cowbane* (*Cicuta virosa*) about $1\frac{1}{4}$ per cent. of an essential oil composed of *Cymol* or *Cymene* ($C_{10}H_{14}$), and *Ouminol* ($C_{10}H_{12}O$). The latter is an aldehyd readily uniting with alkaline bisulphites and by oxidation yielding *Ouminic acid* ($C_{10}H_{12}O_2$). *Cymol* also occurs in *Garden Thyme* (*Thymus vulgaris*). 28. *Eucalyptus globulus* furnishes an oil the more volatile and chief portion of which is *eucalyptol* or *cymene* and a *terpene* ($C_{10}H_{14} + 2C_{10}H_{16}$), together with an oxidized portion, $C_{10}H_{14}O$ and $C_{10}H_{16}O$. 29. *Elecampane-root* (*Inula Helenium*) by distillation with water yields solid volatile *helenin* (C_6H_8O), a camphor-oil or *inulol* ($C_{10}H_{16}O$) and *inulic anhydride* ($C_{15}H_{22}O_3$). 30. Oil of *Fennel-fruit* (*Fœniculi Fructus*, B. P.) differs in odour, but contains the same proximate constituents as oil of anise. 31. Oil of *Geranium*, or *Ginger Grass* oil, from *Andropogon schœnanthus*, and various species of *Pelargonium*, contains *geraniol* ($C_{10}H_{18}O$). 32. *Grains of Paradise* (*Amomum melegueta*), *Guinea Grains* or *Melegueta Pepper*, *Semina Cardamomi Majoris*, contain essential oil ($C_{10}H_{16}$ and $C_{10}H_{16}O$) and a highly pungent resin. 33. Oil of *Juniper* (*Oleum Juniperi*, B. P.) contains a hydrocarbon ($C_{10}H_{16}$) which by contact with water yields a white crystalline hydrous compound ($C_{10}H_{16}H_2O$) and a polymeric hydrocarbon ($C_{20}H_{32}$). 34. Oil of *Lavender* (*Oleum Lavandulæ*, B. P.) has not been satisfactorily examined. 35. Oil or butter or camphor of *Orris* (*Iris florentina*) is a soft solid lighter than water. Flückiger and Hanbury found it to be chiefly myristic acid associated with a little essential oil. 36. Oil of *Peppermint* (*Oleum Mentha Piperitæ*, B. P.) consists of a hydrocarbon, *menthene* ($C_{10}H_{18}$), different from that of most volatile oils, and hydrous *menthene* ($C_{10}H_{18}H_2O$), a crystalline stearopten. 37. Oil of *Spearmint* (*Oleum Mentha Viridis*, B. P.), the Common Mint of the kitchen garden, contains a liquid having the formula $C_{10}H_{20}O$ or $C_{10}H_{18}H_2O$, also, according to Gladstone, *menthol* ($C_{10}H_{14}O$), isomeric with carvol. 38. Oil of *Pennyroyal* (*Mentha pulegium*) contains, according to Kane, $C_{10}H_{16}O$. 39. Oil of *Nutmeg* (*Oleum Myristicæ*, B. P.) is composed of a hydrocarbon (*myristicene* ($C_{10}H_{16}$) and *myristicol* ($C_{10}H_{16}O$)—Gladstone. *Mace*, the arillus or net-like envelope of the nutmeg, appears to yield similar bodies. 40. Oil or *Otto* or *Attar*

of *Cabbage-Rose petals* (*Rosæ Centifoliæ Petala*, B. P.) gives the fragrance to Rose-water (*Aqua Rosæ*, B. P.). It resembles most other volatile oils in being composed of a hydrocarbon and an oxidized portion, but differs from all in this respect, that the hydrocarbon is solid and is destitute of odour, while the oxygenated constituent is liquid and the source of the perfume. According to Flückiger the solid hydrocarbon ($C_{18}H_{16}$) yields succinic acid as the chief product of its oxidation by nitric acid, and in other respects affords evidence of belonging to the paraffin series of fats. 41. Oil of *Rosemary-tops* (*Oleum Rosmarini*, B. P.) exists in the plant to the extent of from $1\frac{1}{2}$ to 3 parts per 1000. It chiefly consists of a hydrocarbon ($C_{10}H_{16}$) resembling that from Myrtle, *Myrtus communis*, but also contains oxygenized oil, and stearopten ($C_{10}H_{16}O$ and $C_{10}H_{18}O$) in variable proportions. 42. Oil of *Rue* (*Oleum Rutæ*, B. P.) contains a small quantity of hydrocarbon ($C_{10}H_{16}$) with some rutiic aldehyd ($C_{10}H_{20}O$), but, according to Greville Williams, is chiefly euodic aldehyd ($C_{11}H_{22}O$), some lauric aldehyd ($C_{12}H_{24}O$) also being present. Gorup-Besanez and Grimm have obtained oil of rue ($C_{11}H_{22}O$) artificially as one of the products of the destructive distillation of acetate and caprate of calcium. 43. Oil of *Savin* (*Oleum Sabinæ*, B. P.) contains several hydrocarbons, but none isomeric with oil of turpentine (Tilden). 44. Oil of *Elder-flowers* (*Sambuci Flores*, B. P.) occurs in very small quantity; it has a butyraceous consistence; it contains a hydrocarbon, *sambucene* ($C_{10}H_{16}$), and probably a camphor. 45. Oil of *Sandal-wood* has not been thoroughly examined. It occurs to the extent of about 1 per cent. in the fragrant white or yellow sandal-wood of India, *Santalum album*, a small tree of the natural order Santalacæ, and not to be confounded with the *Pterocarpus santalinus*, a tree of the natural order Leguminosæ, and furnishing the inodorous Red Sandal-wood or Red Sander's Wood of the dyer. 46. Oil of *Sassafras-root*, sp. gr. 1.094 (*Sassafras Radix*, B. P.), yields *safren* ($C_{10}H_{16}$) and large quantities of a stearopten, *safrol* or *sassafröl* ($C_{10}H_{10}O_2$). 47. Oil of *Mustard* (*Oleum Sinapis*, B. P.) is the sulphocyanate of allyl (p. 532). If adulterated with alcohol, its sp. gr. is below 1.015. 48. Oil of *Sweet Flag* (*Acorus calamus*) contains the hydrocarbon $C_{10}H_{16}$. (The rhizome also contains *Acorin*, a bitter glucoside.) 49. Oil of *Turpentine* (*Oleum Terebinthinæ*, B. P.). Turpentine itself is really an oleo-resin of about the consistence of fresh honey.

It flows naturally or by incision from the wood of most coniferous trees, larch (*Larix Europæa*) yielding *Venice turpentine*, *Abies balsamea* furnishing *Canadian Turpentine* or *Canada Balsam* (*Terebinthina Canadensis*, B. P.), the bark of *Pistachia terebinthus* the variety termed *Chian Turpentine* (containing about 1 part of essential oil to 7 of resin), and the *Pinus palustris*, *P. abies*, *P. pinaster*, and *P. tæda* affording the common, American, Turpentine. *Pinus maritima* gives the French or Bordeaux Turpentine, and *P. picea* the old fragrant Strasburg Turpentine. By distillation turpentine is separated into *rosin* or *resin* (p. 524), which remains in the still, and *essential oil of turpentine*, often termed simply *turpentine*, *spirit of turpentine*, or "*turps*," which distills over. Mixed with alkali to saturate resinous acids, and redistilled, oil of turpentine furnishes about 80 per cent. of *rectified oil of turpentine*. Under the influence of heat, chemical agents, or both, pure oil of turpentine ($C_{10}H_{16}$) yields many derivatives of considerable chemical interest. The terpene of Bordeaux turpentine (terebenthene) rotates a ray of polarized light more than, and in the opposite direction to, the terpene of American turpentine. 50. Oil of common garden *Thyme* (*Thymus vulgaris*) is composed of *cymene* or *cymol* ($C_{10}H_{14}$), *thymene* ($C_{10}H_{16}$), and *thymol* ($C_{10}H_{14}O$). 51. Oil of *Turmeric* (*Curcuma longa*) resembles thymol. Thymol crystallizes out when oil of thyme or of ptychotis, etc., is kept at a low temperature for a day or two. It may also be obtained by shaking the oils with caustic alkali, and treating the separated alkaline liquid by an acid. It may be purified by distillation or by crystallization from alcohol. It would seem that as an antiseptic thymol is far stronger than carbolic acid. 52. Oil of *Valerian-root* (*Valerianæ Radix*, B. P.) is a mixture of a hydrocarbon, *valerene* or *borneene* ($C_{10}H_{16}$), and *valerol* ($C_6H_{10}O$). Valerol slowly oxidizes to valerianic acid, known by its smell. A similar change occurs at once if oil of valerian be allowed to fall, drop by drop, on heated caustic potash: $C_6H_{10}O + 3KHO + H_2O = K_2CO_3 + KC_5H_9O_2 + 3H_2$. By the action of sulphuric acid on the valerianate of potassium thus produced, valerianic acid is obtained. 53. Oil of *Verbena*, *Lemon Grass Oil*, or *Indian Melissa Oil*, is obtained from *Andropogon citratus* (*Oleum Andropogi Citrati*, P. I.). 54. Oil of *Ginger* (*Zingiber*, B. P.) contains a fluid having the composition of hydrous oil of turpentine.

(For an analysis of ginger, by Thresh, and for a paper on

"Soluble Essence of Ginger," see the *Pharmaceutical Journals* for August 30 and September 6, 1879.)

CAMPHORS.—In addition to the stearoptens or camphors already mentioned as being contained in or formed from volatile oils, there is one that is a common article of trade. It is obtained from the wood of *Camphora officinarum*, or Camphor-Laurel, in Japan (termed, in Europe, Dutch camphor because imported by the Dutch) and in China (known as Formosa camphor), by a rough process of distillation with water, and is resublimed in this country (*Camphora*, B. P.). The formula of laurel-camphor is $C_{10}H_{16}O$. Bromine heated with camphor gives *monobrom-camphor* ($C_{10}H_{15}BrO$) and hydrobromic acid. Monobrom-camphor is camphor in each molecule of which an atom of hydrogen has been displaced by one of bromine. Recrystallized, it occurs in white prisms. The essential oil, from which doubtless camphor is derived by oxidation, is easily obtained from the wood, and is occasionally met with in commerce under the name of *liquid camphor* or *camphor-oil*; its formula is $C_{20}H_{32}O$; by exposure to air it becomes oxidized and deposits common camphor, $2C_{20}H_{32}O + O_2 = 4C_{10}H_{16}O$. Camphor distilled with phosphoric anhydride yields cymol, $C_{10}H_{14}$. There is another kind of camphor, in European markets less common than laurel-camphor, but highly esteemed by the Chinese: it is obtained from the *Dryobalanops aromatica*, and denominated Sumatra or Borneo camphor. It differs slightly from laurel-camphor in containing more hydrogen, its formula being $C_{10}H_{18}O$. It is accompanied in the tree by a volatile oil ($C_{10}H_{16}$) isomeric with oil of turpentine. This oil, *borneene*, is also occasionally met with in trade under the name of *liquid camphor* or *camphor-oil*, but differs from laurel-camphor oil in not depositing crystals on exposure to air.

Camphor is soluble to a slight extent in water (40 grains per gallon, Pooley). The official Camphor-water (*Aqua Camphora*, B. P.), or *Camphor mixture*, is such a solution.

Common camphor, and many other of the camphors, oily hydrocarbons, and oxidized hydrocarbons, yield *camphoric acid* ($C_{10}H_{16}O_4$) and *camphoretic acid* ($C_{10}H_{14}O_7$) when attacked by oxidizing agents. Such reactions indicate natural relationships. Camphoric acid is a good antiseptic.

Cantharidin ($C_8H_6O_3$?), the active blistering principle of cantharides (*Cantharides*, B. P.) and other vesicating insects (such as *Myiatriis cichorii* or *Telini Fly*, P. I., common in

India), has most of the properties of a camphor or stearopten. It slowly crystallizes, from an alcoholic tincture of the beetles, in fusible, volatile, micaceous plates. The following process for the extraction of cantharidin is by Fumouze:—Powdered cantharides are macerated with chloroform for twenty-four hours; and this treatment is repeated twice with fresh quantities of solvent, the residue having been well squeezed each time. The collected solutions are then distilled, and the dark green residue treated with bisulphide of carbon, which dissolves fatty, resinous, and other matters, and precipitates the cantharidin. The precipitate is thrown on a filter, washed with bisulphide of carbon, and recrystallized from chloroform. The same process, omitting the final recrystallization, may be used for the quantitative estimation of cantharidin in cantharides. The average quantity found is from four to five, or occasionally ten or even twelve, parts in one thousand. Cantharidin is readily soluble in warm glacial acetic acid (Tichborne) and still more readily in acetic ether. Cantharides from which the fat has been removed by petroleum ether yield their cantharidin with great facility.

Massing and Dragendorff consider cantharidin to be an anhydride ($C_5H_6O_2$), and that with the elements of water it forms *cantharidic acid* ($H_2C_5H_6O_3$). A cantharidate of potassium has the composition $KHC_5H_6O_3$. Piccard gives the vapour density of cantharidin as about 6.5 and its formula $C_{10}H_{12}O_4$.

QUESTIONS AND EXERCISES.

837. How do volatile oils usually differ chemically from fixed oils?
838. What are the general chemical characters of volatile oils?
839. Describe the usual process by which volatile oils are obtained.
840. Mention the differences in composition between the volatile oils of *Anthemis nobilis* and *Matricaria chamomilla*.
841. Give the systematic name of oil of horseradish.
842. State the general composition of the oils of lemon, lime, bergamot, citron, and cedar.
843. Name the constituents of oil of cloves.
844. In what respect does oil (or otto) of roses differ from other volatile oils?

845. To what class of substances do the constituents of oil of rue belong?

846. How does natural turpentine differ from the turpentine of trade?

847. With what object is commercial turpentine rectified?

848. How is camphor-oil related to camphor?

849. In what respects do Borneo or Sumatra camphor and camphor-oil differ from the corresponding products of Japan and China?

850. What is the nature of cantharidin?

RESINOID SUBSTANCES.

RESINS, OLEO-RESINS, GUM-RESINS, BALSAMS.

Resins occur in plants generally in association with volatile oils. They closely resemble camphors or stearoptens, but are not volatile, and differ from oil and fats mainly in being solid and brittle. *Oleo-resins* are mixtures of a resin and a volatile oil. *Gum-resins* are mixtures of a resin or oleo-resin and gum. *Balsams* are commonly described as resins or oleo-resins which yield benzoic or cinnamic acid; but oleo-resins containing neither of these acids are often termed balsams, e.g. balsam of copaiva and Canada balsam. A physico-chemical method for the identification of the chief resins, gum-resins, and balsams will be found in the *Pharmaceutical Journal* for November 17, 1877.

Resins appear to be somewhat antiseptic. Beer is said never to turn sour in casks lined with Burgundy pitch. The resin of hops has, perhaps, a similar effect in retarding oxidation of alcohol.

RESINS.—1. *Resin*, *rosin*, or *colophony* (*Resina*, B. P.) is the type of this class. Its source is the oleo-resin or true turpentine of the conifers, a body which by distillation yields spirit of turpentine and a residuum of rosin. "Brown" and "White" rosin are met with in trade. The former is the residue of American, the latter of Bordeaux turpentine (from *Pinus Abies*, etc., and *Pinus Maritima* respectively). The chief constituents of brown resin are *pinic acid* ($\text{HC}_{20}\text{H}_{29}\text{O}_2$) and *sylic acid*, identical in composition, but differing in properties (*vide* Isomerism), the former being soluble and the latter insoluble in cold spirit of wine. White resin or "gali-

pot" is chiefly *pimaric acid*, also isomeric with *pinic acid*. *Pinic acid* cautiously heated yields *colophonic* or *colopholic acid*. Among the products of the destructive distillation of resin, Tichborne has found "*colophonic hydrate*" ($C_{10}H_{22}O_3 \cdot H_2O$), a white inodorous crystalline substance, and by depriving this of water has obtained white crystalline *colophonins* ($C_{10}H_{22}O_3$). Resin is soluble in oil of turpentine. Contact with sulphuric acid immediately colours it strongly red. It is a constituent of eight of the fourteen Plasters (*Emplastra*) of the British Pharmacopœia. 2. *Arnicin* ($C_{30}H_{50}O_4$), the chief acrid if not the only active principle of *Arnica-root* (*Arnica Radix*, B. P.), is a resin. 3. *Cannabin*, said to be the active principle of *Indian Hemp* (*Cannabis Indica*, B. P.), was obtained in 1846 by T. and H. Smith, and is a resin. Personne in 1857 isolated a volatile oil, also said to possess much medicinal activity, consisting of *cannabene* ($C_{18}H_{30}$) and a solid crystalline "*hydride of cannabene*" ($C_{18}H_{22}$). Preobraschensky states that the active principle is nicotine. 4. *Capiscum-fruit* contains a resin (p. 562). 5. *Castorin*, a resinous matter, is the name given to the chief constituent of *Castor* (*Castoreum*, B. P.), the dried preputial follicles and included secretion of the Beaver (*Castor Fiber*). 6. *Copal*.—The best copal is the exuded resin of trees of extinct forests, and is found beneath the surface of the ground in the neighbourhood of existing trees. It appears to be a mixture of acids, but its character is still obscure. 7. *Dragon's Blood*, a crimson-red resin found as an exudation on the mature fruits of a Rotang or Rattan Palm (*Calamus draco*). It is said to be a definite substance, and to have the formula $C_{20}H_{20}O_4$. 8. *Ergotin* is a very active resinoid constituent of *Ergot* (*Ergota*, B. P.), or "the sclerotium (compact mycelium or spawn) of *Claviceps purpurea*, produced within the paleæ of the common rye, *Secale cereale*." According to Wenzell, ergot contains two alkaloids, *ecboline* and *ergotine*, to the former of which, he says, the activity of ergot is due. Blumberg considers these alkaloids to be identical. Tanret states that an unstable alkaloid termed *ergotinine* occurs in ergot to the extent of 1 per 1000, and that it is accompanied by a camphor. Dragendorff and Podwissotzky assert that ergot owes most of its activity to *sclerotic* or *sclerotinic acid*, present to the extent of about 4 per cent. No separated principle representing the full activity of ergot has yet been extracted from that drug. 9. *Guaiacum-resin* is a mixture of several substances (p. 496). 10. *Jalap-resin* (p. 497). 11.

Kousso (*Cusso*, B. P.) yields yellow crystals of a resinoid body readily soluble in alkaline liquids, *kosin* or *koussein* ($C_{51}H_{38}O_{10}$). 12. *Mastic* (*Mastiche*, B. P.) is a resinous exudation obtained by incision from the stem of the Mastic or Lentisk tree. Nine-tenths of mastic is *mastichic acid* ($C_{30}H_{31}O_2$), a resin soluble in alcohol; the remaining tenth, *masticin* ($C_{30}H_{31}O$), a tenacious elastic resin. 13. *Mezereon*, the dried bark (*Mezerei Cortex*, B. P.) of *Daphne mezereum*, *Mezereon*, and *Daphne laureola*, Spurge Laurel, owes its acridity to a resin. 14. *Pepper* contains a resin (p. 469). 15. *Burgundy pitch* (*Pix Burgundica*, B. P.) is the melted and strained exudation from the stem of the Spruce Fir, *Abies Excelsa*. The term *Burgundy* is a misnomer, the resin never having been collected at or near Burgundy—Finland, and to a smaller extent Baden, and Austria being the countries whence it is derived. Its constituents closely resemble those of common resin. It is often adulterated and imitated by a mixture of resin with palm-oil, water, etc., from which it may be readily distinguished by its duller yellow colour, highly aromatic odour, greater solubility in alcohol, and almost complete solubility in twice its weight of glacial acetic acid (Hanbury). 16. *Podophyllum-resin*.—In preparing the resin of *podophyllum*, or *May-apple* (*Podophylli Resina*, B. P.), an alcoholic extract of the root (*Podophylli Radix*, B. P.) is poured into cold water acidulated by hydrochloric acid. This resin is the chief active principle of *podophyllum-root*. According to Guareschi, *podophyllin* is a mixture of a resin proper and a glucoside resembling convolvulin. 17. *Pyrethrin* is the name of the acrid resinous active principle of the root of *Anacyclus pyrethrum* or *Pellitory-root* (*Pyrethri Radix*, B. P.). According to Buckheim, the action of alkalies breaks it up into piperidine and pyrethric acid. The crystalline poisonous principle obtained by Bellesme from *Pyrethrum carneum*, the powder of which (and of *P. roseum*, and especially *P. cinerariæ folium* or Dalmatian Insect Powder) is the well-known "insecticide," has not yet been analysed. 18. The resins of *Rhubarb* have already been alluded to in connection with Chrysophanic acid (p. 394). 19. *Rottlerin* is the name given by Anderson to a crystalline resin from *Kamala* (*Kamala*, B. P.), the minute glands that cover the capsules of *Rottlera tinctoria*: to this and, apparently, allied resins *Kamala* owes its activity as an anthelmintic.

OLEO-RESINS.—1. *Capsicin*, a term suggestive of a definite

chemical substance, is a name somewhat unhappily accorded to an indefinite substance, an oleo-resin, obtained by digesting the alcoholic extract of Capsicum fruit (*Capsici Fructus*, B. P.) in ether and evaporating the clear ethereal fluid to dryness. Besides volatile oil and resin, capsicum fruits contain much fatty matter which Thresh states is chiefly free palmitic acid. (See also *Capsicine* and *Capsaicin*, p. 466.) 2. *Copaiva* (*Copaiba*, B. P.) is a mixture of essential oil ($C_{15}H_{34}$), with 2 or more per cent. of brown soft resin, and 30 to 60 of a yellow dark crystalline resin termed *Copaivic acid* ($C_{10}H_{30}O_3$). *Copaiva*, containing about equal parts of this acid and of the oil, heated with a fourth of its weight of the official carbonate of magnesium, yields a transparent fluid, owing to the formation of copaiate of magnesium and solution of this soap in the essential oil. With an equal weight of the carbonate enough soap is produced to take up the whole of the essential oil, and form a mass capable of being rolled into pills. A much smaller quantity of calcined magnesia, as might be expected, effects the same result; but more time, often several days, is required before complete reaction is effected. Quicklime has a similar effect. Perhaps carbonate reacts more quickly because of its fine state of division and admixture of hydrate—in which case hydrates of calcium and magnesium may be expected to act better than the calcined preparations, and in much smaller quantity than carbonate of magnesium. *Copaiva*, unlike, 2, *Wood-oil*, or *Gurjun Balsam* (*Dipterocarpi Balsamum*, P. I.), a similar oleo-resin from the *Dipterocarpus turbinatus* (*D. Lævis*, P. I.), does not become gelatinous when heated to 270° F. *Copaiva* is often slightly fluorescent; *Gurjun balsam* is highly fluorescent. The stated analogy of *Gurjun balsam* to *copaiva* is borne out by its chemical composition; for by distillation it yields about 40 per cent. of an essential oil identical in composition with oil of *copaiva*, the non-volatile portion being resinous. The adulteration of *copaiva* with fixed oil is best detected by heating 20 or 30 drops in a capsule until all essential oil has evaporated. (Turpentine is betrayed by its odour during this evaporation.) The residue, *copaiva resin*, is brittle if pure, and more or less sticky or soft if fixed oil is present. The limit of brittleness is stated, by Siebold, to be reached when one per cent. of oil has been added to the *copaiva*, that amount preventing the residue being reduced to a fine powder. 3. *Elemi* (*Elemi*, B. P.) is an exudation from a tree growing in the Philippine Islands.

It consists of volatile oil ($C_{10}H_{16}$) with 80 or more per cent. of two resins, the one ($C_{30}H_{32}O_2$) soluble in cold alcohol, the other *Amyrin* (C_5H_8) $_5$ H_2O almost insoluble, associated with *Amyric acid* (C_5H_8) $_7$ O_4 (Buri). It also contains small quantities of two crystalline bodies soluble in water, *Bryoidin* (C_5H_8) $_4$ H_2O and *Breidin* (Flückiger). The *Icacin* of Stenhouse and Groves is either identical with amyryn or perhaps has the formula (C_5H_8) $_9$ H_2O . All these bodies are probably hydrous terpenes. 4. *Wood-tar* (*Pix Liquida*, B. P.) is a mixture of several resinoid and oily bodies (amongst others Creasote, p. 529) obtained by destructive distillation from the wood of *Pinus sylvestris* and other pines. When heated it yields a terebinthinate oil and a residue of *pitch*. 5. *Turpentine*. These oleo-resins have been mentioned in connection with oil of turpentine, their volatile, and resin, their fixed constituent. 6. *Common Frankincense* (*Thus Americanum*, B. P.) is the concrete turpentine of *Pinus tæda*. 7. *Canada Balsam* (*Terebinthina Canadensis*, B. P.) is largely gathered in the province of Quebec, and is the turpentine or oleo-resin of the Balm of Gilead Fir (*Abies balsamea*). 8. *Sumbul-root* (*Sumbul Radix*, B. P.) seems to owe its stimulating property to two oleo-resins, one soluble in ether, the other in alcohol. 9. *Oleo-resin of Lupulin* (U. S. P.) is an ethereal extract of the yellow powder (*Lupulin*, U. S. P.) attached to the small nuts at the base of the scales which form the aggregate fruit of the Hop (*Humulus Lupulus*). It contains essential oil of hop (valerol, $C_6H_{10}O$), oxidized oil or resin, bitter extract containing the hop-bitter, lupulinic acid ($C_{33}H_{50}O_7$), and tannic acid.

GUM-RESINS.—1. *Ammoniacum* (*Ammoniacum*, B. P.) is an exudation from the *Dorema Ammoniacum*. It contains nearly 20 per cent. of gum, a little volatile oil, and about 70 of resin ($C_{40}H_{50}O_9$ —Johnston). 2. *Asafoetida* (*Asafoetida*, B. P.) is a gum-resin obtained, by incision, from the living root of *Narthez asa-foetida*. It contains from 50 to 70 per cent. of a resin which is partly *ferulaic acid* ($C_{10}H_{10}O_4$), 25 to 30 per cent. of gum (about two-thirds arabin, one-third bassorin, p. 123), and 3 to 5 per cent. of volatile oil. 3. *Euphorbium*, an old drug which is an emetic and purgative resin. It contains an amorphous active resin ($C_{20}H_{32}O_4$), crystalline *euphorbon* ($C_{28}H_{44}O_2$), and mucilage (Flückiger). 4. The ordinary or Siam *Gamboge* (*Cambogia*, B. P.) of European trade is obtained from the *Garcinia morella*; the gamboge of

India (*Cambogia Indica vel Mysoriensis*, P. I.) from *G. pictoria*. When of best quality it contains about 20 per cent. of a gum, and 80 to 75 per cent. of a yellow resin termed *gambogic acid* ($C_{20}H_{23}O_4$). 5. *Galbanum* (*Galbanum*, B. P.) contains from 20 to 25 per cent. of gum, about 65 per cent. of resin ($C_{40}H_{54}O_7$), and 3 or 4 per cent. of volatile oil. Galbanum heated for some time to 212° F. with hydrochloric acid, the liquid separated and shaken with ether or chloroform, and the latter evaporated, yields somewhat less than 1 per cent. of colourless acicular crystals of *umbelliferone* ($C_9H_6O_3$)*. 6. *Myrrh* (*Myrrha*, B. P.), an exudation from the stem of *Balsamodendron myrrha*, contains about half its weight of soluble arabinoid gum, 10 per cent. of insoluble gum (probably bassorin), $2\frac{1}{2}$ of volatile oil, and about 25 per cent. of resin (myrrhic acid). (For a note by R. H. Parker on the spurious Gums imported with Myrrh see the *Pharmaceutical Journal* for July 17, 1880.) 7. *Olibanum* (P. I.), *Thu* *masculum* or *Arabian Frankincense* (from various species of *Boswellia*), is about one-third gum and nearly two-thirds resin ($C_{40}H_{30}O_6$) with a little hydrocarbon ($C_{10}H_{16}$) and oxidized hydrocarbon volatile oils. It has always been an important ingredient of *incense*—myrrh, storax, benzoin, and such fragrant combustible resinous substances being other constituents. 8. *Scammony* (p. 500).

Gum-resins need only to be finely powdered and rubbed in a mortar with water to yield a medicinal *emulsion*, in which the fine particles of resin are held in suspension by the aqueous solution of gum.

BALSAMS.—1. *Benzoin* (*Benzoinum*, B. P.) is obtained from

* "Umbelliferone is soluble in water; its solution exhibits, especially on addition of an alkali, a brilliant blue fluorescence which is destroyed by an acid. If a small fragment of galbanum is immersed in water no fluorescence is observed, but it is immediately produced by a drop of ammonia. The same phenomenon takes place with *asafoetida*, and in a slight degree with *ammoniacum*; it is probably due to traces of umbelliferone preexisting in those drugs. Umbelliferone is also produced from many other aromatic umbelliferous plants, as *Angelica*, *Levisticum*, and *Meum*, when their respective resins are submitted to dry distillation; also from the resin of *Daphne mezereum*. The fluorescence of umbelliferone may be beautifully shown by dipping some bibulous paper into water which has stood for an hour or two on lumps of galbanum, and drying it. A strip of this paper placed in a test-tube of water with a drop of ammonia will give a superb blue solution, instantly losing its colour on the addition of a drop of hydrochloric acid." (Flückiger and Hanbury.)

incisions in the bark of *Styrax benzoin*. It contains from 12 to 15 per cent. of benzoic acid (p. 392), about 50 per cent. of a resin (α) soluble in ether, 25 to 30 per cent. of a resin (β) soluble in alcohol only, and 3 to 4 per cent. of a resin (γ) soluble in solution of carbonate of sodium. The α resin is considered to be a compound of the β ($C_{40}H_{46}O_9$) and the γ ($C_{30}H_{40}O_5$). The balsams of Peru, Tolu, and Storax differ from benzoin in containing cinnamic (p. 394) in place of benzoic acid; hence they yield, by oxidation, hydride of benzoyl (oil of bitter almonds).

2. *Balsam of Peru* (*Balsamum Peruvianum*, B. P.), from the *Myroxylon Pereiræ*, is a mixture of oily matter with about one-quarter or one-third resinous matter, and 6 per cent. of cinnamic acid. The oil, by fractional distillation in an atmosphere of carbonic acid gas and under diminished pressure, furnishes benzylic alcohol (C_7H_7HO), benzoate of benzyl ($C_7H_7C_7H_5O_2$), and cinnamate of benzyl ($C_7H_7C_9H_7O_2$) or cinnaméin (Kraut). By action of alcoholic solution of potash it yields benzoate and cinnamate of potassium, and benzylic alcohol; also cinnamic alcohol (C_9H_9HO), otherwise known as *peruvine* or *styrone*; it also often holds in solution *metacinnaméin* or *styracin* ($C_{18}H_{16}O_3$), isomeric with hydride of cinnamyl (C_9H_7OH). The resin of Peru seems to result from the action of moisture on the oil.

3. *Balsam of Tolu* (*Balsamum Tolutanum*) is an exudation from incisions in the bark of *Myroxylon toluifera*; in composition it closely resembles balsam of Peru, but is more susceptible of resinification. It contains benzoate and cinnamate of benzyl, cinnamic acid, a little benzoic acid (Busse) and about 1 per cent. of a volatile hydrocarbon, *tolene*, $C_{10}H_{16}$. Old hard balsam of tolu is a convenient source of cinnamic acid, which may be extracted by the same process as that by which benzoic acid is obtained from benzoin—namely, ebullition with alkali, filtration, and precipitation by hydrochloric acid.

4. *Storax* is an oleo-resin obtained from the *Liquidambar orientale*. It contains a volatile oil termed *styröl*, *cinnamene*, or *cinnamol* (C_8H_8),—which possibly (Berthelot) is condensed acetylene, $4C_2H_2$ —cinnamic acid, styracin, or cinnamate of cinnamyl ($C_9H_9C_9H_7O_2$), and a soft and a hard resin. Styröl differs from similar hydrocarbons in being converted into a polymeric solid, termed *metastyrol* or *draconyl*, on heating to about 400° F. For medicinal use, storax (*Styrax Præparatus*, B. P.) is purified by solution in alcohol, filtration, and removal of the alcohol by distillation.

Caoutchouc or India-rubber, and Gutta Percha.

Caoutchouc is the hardened juice of *Hevea* (*Siphonia*) *Brasiliensis*, *Ocstilloa elastica*, *Urceola elastica*, *Ficus elastica*, and other plants (Collins). Heated moderately with sulphur it takes up 2 or 3 per cent., and forms *vulcanised india-rubber*; at a higher temperature a hard horny product, termed *ebonite* or *vulcanite*, results. *Gutta Percha* (*Gutta Percha*, B. P.) is the concrete drop or juice of the *percha* (Malay) tree, the *Isonandra gutta*, and of other Sapotaceous plants. White gutta percha is obtained by precipitating a solution of ordinary gutta percha in chloroform by alcohol, washing the precipitate with alcohol, and finally boiling in water and moulding into the desired form while still hot. The official solution of Gutta Percha (*Liquor Gutta Percha*, B. P.) is made by digesting *thin* slices of gutta percha in 12 parts by weight of chloroform, and then "fining" by shaking with 1 part of carbonate of lead and setting aside till the fluid is clear.

These two elastic substances, in the pure state, are hydrocarbons (xC_6H_8), usually slightly oxidized.

QUESTIONS AND EXERCISES.

851. How do resins occur in nature? Distinguish between resins and camphors. Mention the points of difference of resins, oleo-resins, gum-resins, and balsams.

852. Name the source and the chief constituents of common Resin or Rosin.

853. Enumerate some official articles of which the active constituents are resins.

854. Give the chief distinguishing characters of Burgundy Pitch.

855. What is the average proportion of oil and of resin in the so-called Balsam of Copaiva?

856. Explain the effect of carbonate of magnesium, magnesia, and lime on copaiva.

857. State the nature of Wood-Tar.

858. Why do Ammoniacum, Asafoetida, Gamboge, Galbanum, Myrrh, and similar substances give an emulsion by mere trituration with water?

859. In what respects does Benzoin differ from the balsams of Peru, Tolu, and Storax?

860. What is the chemical nature of India-rubber and Gutta Percha?

861. How is India-rubber *vulcanized* and converted into *ebonite* or *vulcanite*?

COLOURING-MATTERS.

The animal, vegetable, and mineral kingdoms abound in substances or pigments which powerfully decompose light, absorbing certain of its constituent colours, and reflecting some others. Thus, for example, most leaves contain a body termed chlorophyl, which has the property of absorbing red light and reflecting green; these reflected rays entering the eye of an observer, and striking on the retina (the expanded extremity of the optic nerve), always communicate the same impression to the brain; in popular language the leaf is said to be green. Art has richly supplemented the number of such natural colouring matters.

YELLOW.—1. *Chrome-yellow* occurs in more than a dozen shades (see Lead, chromate of). 2. *Fustic* or *yellow wood* is the wood of the *Rhus cotinus*. 3. *Gamboge* (see Gamboge). 4. *Ochre* is met with of many tints, under the names of *yellow ochre*, *gold yellow*, *gold earth* or *ochre*, *yellow sienna*, *Chinese yellow*. It is chiefly a mixture of oxyhydrates of iron with alumina and lime. 5. *Orpiment* is a sulphide of arsenicum (As_2S_3). 6. *Persian berries* or *Avignon grains* contain a yellow principle termed *rhamnin* and other crystalline bodies: they are the product of the *Rhamnus infectorius*. 7. *Purree* or *Indian yellow* is said by Stenhouse to owe its colour to *purrate* or *euzanthate* of magnesium ($\text{MgC}_{42}\text{H}_{34}\text{O}_{22}$). 8. *Quercitron* is the bark of *Quercus tinctoria*: it contains the yellow glucoside, *quercitrin* ($\text{C}_{18}\text{H}_{18}\text{O}_{10}$, H_2O). 9. *Rhubarb* (see Chrysophanic acid, p. 394). 10. *Saffron* (*Crocus*, B. P.), the dried stigma and part of the style of *Crocus sativus*, yields *saffranin* or *polychroite*, an orange-red glucoside, which by the action of dilute acids and by other means breaks up as shown in the following equation, yielding red *crocin* (Weiss).



11. *Turmeric*, the rhizome of *Curcuma longa*, owes its yellow

colour to *curcumin*, a resinous matter, the formula of which is said by Daube to be $C_{10}H_{10}O_3$, and by Iwanof C_2H_4O . Possibly two yellow pigments are present. 12. *Weld* (*Reseda luteola*) contains a durable yellow matter termed *luteolin* ($C_{20}H_{14}O_3$). 13. *Picric* or *carbazotic acid* (p. 530) is a very powerful yellow dye. 14. Dried and powdered carrots yield to bisulphide of carbon a yellow colouring-matter, "carrotin," which is obtained on evaporating the solvent. It is said to be used in colouring butter.

RED.—1. *Alkanet*, the root of *Alkanna tinctoria*, Tausch, *Anchusa tinctoria*, Desf., yields *anchusin* ($C_{35}H_{40}O_3$), a resinoid matter soluble in oils and fat. 2. *Annatto*, *Arnatto*, or *Arnotto*, a paste prepared from the seeds of *Bixa orellana*, contains *bixin*, an orange-red, and *orellin*, a yellow principle. 3. *Brazil-wood* (*Cæsalpinia brasiliensis*) furnishes *brazilin*, the basis of several lakes. *Sapan-wood* and *Cam-wood* probably contain the same substance. 4. *Cinnabar*, Chinese red, Vermilion, or Paris red, is mercuric sulphide. 5. *Chrome-red* is an oxychromate of lead. 6. *Cochineal* (p. 394). 7. *Madder*, the root of *Rubia tinctorum*, powdered and treated with sulphuric acid and acidulated water to effect the removal of earthy and other inert matters, furnishes a residual powder termed *garancin*. *Garancin* yields to pure water *alizarin* ($C_{14}H_{10}O_4$, $3H_2O$), the red, neutral, crystallizable colouring-matter of madder. *Alizarin* does not exist ready formed in the plant, but is derived, by fermentation, from *rubian*, a yellowish resinoid substance. *Alizarin* is now produced artificially from *anthracene*, one of the solid constituents of coal-tar. 8. *Mulberry-juice* (*Mori Succus*, B. P.) contains a violet-red colouring-matter which has not been chemically examined. 9. *Red lead* (p. 240). 10. *Red oxide of iron*, of shades varying from light to brown red, is found native. The common names of it are Armenian bole, Berlin red, colcothar, English red, red ochre, burnt ochre, red earth, terra di sienna, mineral purple, stone red, and Indian red. 11. *Red Sanders-wood* or *Red Sandal-wood* (*Pterocarpus Lignum*, B. P.), the billets and chips of *Pterocarpus santalinus*, owes its colour to *santalin* ($C_{14}H_{13}O_4$), a crystalline resinoid matter. 12. *Red-Poppy Petals* (*Rhæados Petala*, B. P.), from the *Papaver rhæas*, contains a red colouring principle which has not yet been isolated in a state of purity. The author has sought for morphia in large quantities of the petals, but could not find a trace of that alkaloid. 13. *Red-Rose Petals* (*Rosæ Gallicæ Petala*, B. P.),

GREEN.—1. *Cupro-arseniocal* green pigments (p. 197). 2. *Chlorophyl*, *Leaf-green*, or *Chromule*. A method of extracting chlorophyl is given under "Extracts" (*vide* Index). It is resinoid, soluble in alcohol and ether, insoluble in water, and, according to Frémy, consists of a blue substance, *phyllocyanin* ($C_{34}H_{48}N_4O_{17}$?), and a yellow, *phylloxanthin*; the yellow tints in fading autumnal leaves, he says, are due to the latter principle, the former being the first to fade. Chlorophyl would probably well repay extended investigation. 3. *Sap-green*, *buckthorn*-, *vegetable*-, or *bladder-green* is obtained by evaporating to dryness a mixture of lime and the juice (*Rhamni Succus*, B. P.) of the berries of the Buckthorn (*Rhamnus catharticus*). It is soluble in water, slightly in alcohol, and insoluble in ether and oils. 4. *Green ultramarine* is made by a process similar to that for blue ultramarine. 5. Mixtures of blue and yellow pigments and dyes are common sources of green colours. 6. Glass and earthenware are coloured green by oxide of chromium and black oxide of copper.

BROWN.—1. *Umber*, *Sienna*, or *Chestnut-brown* is found native. By heat it is darkened in tint, and is then known as *burnt umber*. It is a mixture of oxide of iron, silica, and alumina. 2. *Sepia* is a dried fluid from the ink-bag of cuttle-fishes (*Sepiade*); by its ejection into adjacent water the animal is said to obtain opportunity of escape from enemies. 3. *Catechu* (p. 419) furnishes a brown colouring-matter.

BLACK.—1. *Blacklead* (p. 22), *bone-black* (p. 120), or *ivory-black* and *lampblack*, the latter a deposited soot from the incomplete combustion of resin and tar, are varieties of carbon. 2. *Burnt sugar* or *caramel* (p. 489). 3. *Indian ink* is usually a dried mixture of fine lampblack and size or thin glue. 4. *Black ink* is essentially tannates and gallates of iron suspended in water containing a little gum in solution. 5. *Printer's ink* is well boiled linseed or other oil, mixed with good lampblack, vermilion, or other pigment. 6. *Black dyes* are of the same nature as ink. 7. The *pigmentum nigrum* of black feathers, such as those of the common rook, of dark hair, and probably also of the skin of the negro, seems to be due to the black substance which remains undissolved when black feathers are digested for some time in dilute sulphuric acid. It is said to have the formula $C_{18}H_{16}N_2O_8$. (Hodgkinson and Sorby).

WHITE PIGMENTS.—1. *Chalk* or *Whiting* (p. 118). 2. *French chalk*, *steatite*, or *soapstone*, a silicate of magnesium.

3. *Heavy white* (p. 110). 4. *Pearl-white* (p. 289). 5. *Plaster of Paris* (p. 112). 6. *Starch* (p. 473). 7. *White lead* (p. 238). 8. *Zinc white* (p. 145). 9. Oxides of tin and zinc and phosphate of calcium are employed for giving a white opacity to glass.

ANILINE COLOURS. *Coal-tar colours.*—Within the last ten years nearly every shade of colour seen in the animal and vegetable kingdom has been successfully imitated by certain dyes and pigments primarily derived from a mineral, coal. Coal distilled for gas furnishes tar or gas-tar. Coal-tar contains some aniline; but especially it contains a liquid convertible into aniline, namely benzol (C_6H_6), first discovered by Faraday in compressed oil-gas. From aniline, by oxidation, Runge obtained the violet-colour reaction, the body producing which Perkin afterwards studied and isolated, and manufactured under the name of *mauve*. *Aniline-red* (*fuchsine*, *magenta* or *roseaniline*), *aniline-yellow*, *aniline-green*, *aniline-blue*, and, in short, aniline-dyes, lakes, and pigments of every hue of the rainbow, are now common articles of trade. Their application has revolutionized the arts of the dyer and colour-printer.

QUESTIONS AND EXERCISES.

862. Explain the production of colour by the various natural and artificial pigments.

863. Mention the chief yellow colouring-matters, and describe their chemical nature.

864. What is annatto?

865. Name the colorific constituent of madder. Can it be made artificially?

866. State the source of Litmus.

867. Distinguish between Prussian blue and Turnbull's blue; and state how they are manufactured.

868. How is blue ultramarine obtained? How is it affected by acids?

869. Describe the chemical nature of the colouring principle of green leaves.

870. By what agents is glass coloured green?

871. Whence is sepia obtained?

872. Describe the chemistry of black ink.

873. Write a few sentences on *aniline colours*.

QUALITATIVE ANALYSIS OF SUBSTANCES HAVING UNKNOWN PROPERTIES.

Substances are presented to the analyst in one of the three forms in which all matter exists—namely, solid, liquid, or gaseous; and they may contain animal or vegetable as well as mineral matter.

The method of analysis in the case of *solid mineral bodies* has been described on pp. 433 to 442.

Solid animal or vegetable substances (or mixtures of these with mineral bodies) may be indefinite and beyond the grasp of chemistry, or definite and quite within the range of proximate qualitative organic analysis. The presence of such substances is indicated in the preliminary examination of a solid (pp. 434 to 436) by charring and other characters. If no charring occurs, and no volatile liquid is expelled by heat, the absence of such matter is indicated. But if organic matter is present, an endeavour is made to ascertain its precise character. The analyst's knowledge of the history of the substance, or the circumstances under which it comes into his hands, will probably afford a clue to its nature, and enable him to search directly for its proximate constituents. If no such information is at hand, the action of solvents may be employed as likely to afford indication of the general, if not of the precise, nature of the substance. Water, alcohol, ether, chloroform, bisulphide of carbon, each both hot and cold, may in turn be agitated with the substance, the mixture filtered, a portion of the filtrate evaporated, at first partially, setting the product aside, and afterwards to dryness, and any deposit or residue examined with and without the aid of a microscope. Other portions of the filtrate may be treated with acids, alkalies, and solutions of such metallic salts as are commonly used as group-tests for acidulous radicals (p. 431). The action of alkalies, as well as acids weak and strong, hot and cold, may also be tried on the solid substance itself, and colours, odours, and, in short, any effect whatever duly noted. A portion of the substance should also be burnt in an open porcelain crucible until no carbon remains, and the ash, if any, examined; its amount and nature may afford information leading to the identification of the substance.

The foregoing experiments having been carefully performed,

and all results entered in the note-book, a little reflection will possibly lead to the recognition, or may suggest further direct experiments or confirmatory tests, or will, at least, have pointed to the absence of 90 or 95 per cent. of all possible substances, and thus have restricted the area of inquiry to narrow limits. The success attainable in qualitative proximate organic analysis by the medical or pharmaceutical student will of course largely depend on the thoroughness with which the operator has prosecuted his study of practical chemistry generally; but it also will be considerably affected by the extent to which he has cultivated the art of observation, and the opportunities he has had of acquiring a knowledge of the appearance, uses, and common properties of definite chemical substances, and of articles of food, drink, and medicine. The most successful of several good analysts will be the one who has most common sense and most experience.

The pharmaceutical student, who has probably already had some years of experience in pharmacy, occupies an unusually favourable position for prosecuting the proximate analysis of organic and inorganic substances, or, at all events, of that large proportion of such bodies met with in the domain of hygiene and pharmacy. Many substances he will identify at sight, or by aid of a lens, or after applying some simple physical or chemical test. Nor should he find much difficulty, after reaching the present point of practical study, in deciding whether the solid substance under examination belongs to the class of organic acids, organic salts of metallic radicals, alkaloids, salts of alkaloids, amylaceous matter, gums, saccharine substances, glucosides, albumenoid matters, fats, soaps, resins, colouring-matters, etc. For instance, the pharmaceutical student will find less difficulty than the general student in successfully analysing a substance occurring in "scales," because he has experience of the appearances of compounds commonly produced in that form, and because, even if the appearance is new to him, he knows what kind of substances most readily lend themselves to production in that form. While the general student is testing generally, and proceeding cautiously, or searching for general information in books of reference, the pharmaceutical or medical student has incinerated some of the material, noticed whether or not the ash is red (iron) and strongly alkaline (potassium), treated more of the material with an alkali (for ammonium), added excess of ammonia, and examined the precipitate (for cinchonia

or quinia), or shaken up the alkaline liquid successively with ether and chloroform, and tested the residue of these decanted and evaporated solvents (quinia, beberia, strychnia), and examined the aqueous solution of the material, or one of the filtered alkaline liquids, in the usual way for acidulous radicals (citric, tartaric, sulphuric, hypophosphorous). Or he has modified his methods to include search for some "scale preparation" which his special knowledge tells him has been newly introduced to, or is rare in, pharmacy.

In the case of liquids, the solvents as well as the dissolved matters claim attention. A few drops are evaporated to dryness on platinum foil to ascertain if solid matter of any kind is present; the liquid is tested by red and blue litmus-paper to ascertain if free alkalies, free acids, or neither are present; a few drops are heated in a test-tube and the odour of any vapour noticed, a piece of glass tubing bent to a right angle being, if necessary, adapted to the test-tube by a cork, and some of the distilled liquid collected and examined; finally, the usual group-reagents for the several basylous and acidulous radicals are consecutively applied.

Proceeding in this way the student who has already had some experience in pharmacy will not be likely to overlook such solvents as water, acids, alkalies, alcohol, glycerine, ether, chloroform, benzol, fixed oils, and essential oils, or to mix the substances which these menstrua may hold in solution. He will probably also recognise such liquids as carbolic acid, formic acid, lactic acid, methylic alcohol, aldehyd, aniline, nitrobenzol. He must not, however, suppose that he will always be able to qualitatively analyse, say, a bottle of medicine; for the various infusions, decoctions, tinctures, wines, syrups, liniments, confections, extracts, pill-masses, and powders contain vegetable matters most of which at present are quite beyond the reach of the analyst. Neither the highest skill in analysis nor the largest amount of experience concerning the odour, appearance, taste, and uses of drugs is sufficient for the detection of all these vegetable matters. Skill and experience combined, however, will do much, and in most cases even so difficult a task as the one just mentioned be accomplished with reasonable success. Obviously, qualitative analysis alone will not enable the experimenter to produce a mixture of substances similar to that analysed; to this end recourse must be had to quantitative analysis, a subject reserved for subsequent consideration.

Natural fluids, as "Milk" and "Urine" (*vide* Index), admit of special analytical treatment.

Gas-analysis, or *Eudiometry* (from *εἰδία*, *eudia*, calm air, and *μέτρον*, *metron*, a measure, in allusion to the *eudiometer*, an instrument used in measuring the proportion and, as the early chemists thought, the salubrity of the gases of the air), is a branch of experimental investigation, chiefly of a quantitative character, concerning which information must be sought in other treatises. The analysis of atmospheric air from various localities, coal-gas, and gases obtained in chemical researches, involves operations which are scarcely within the sphere of Chemistry applied to Medicine. Beyond the recognition, therefore, of oxygen, hydrogen, nitrogen, carbonic, sulphurous, and hydrosulphuric acid gases, the experimental considerations of the chemistry of gaseous bodies may be omitted. Their study, however, should not be neglected, as existing conceptions of the constitution of chemical substances are largely dependent on the observed relations of the volumes of gaseous compounds to their elements. See previous paragraphs, pp. 37 to 41, 50 to 53, 140 and 141. The best single work on this latter part of the subject is a small book by Hofmann, "Introduction to Modern Chemistry."

Spectral Analysis.—It may be as well to state here that the preliminary and final examinations of minute quantities of solid matter may, in certain cases, profitably include their exposure to a temperature at which they emit light, the flame being physically analysed by a spectroscope. A spectroscope consists essentially of a prism to decompose a ray of light into its constituent colours, with tubes and lenses to collect and transmit the ray or rays to the eye of an observer. The material to be examined is placed on the end of a platinum wire, which is then brought within the edge of a spirit-lamp or other smokeless flame; volatilization, attended usually in the case of a compound by decomposition, at once occurs, and the whole flame is tinged with a characteristic hue. A flat ribbon of rays is next cut off by bringing near to the flame a brass tube, the cap of which is pierced by a narrow slit. At the other end of the tube, at focal distance for parallel rays, is a lens, through which the ribbon of light passes to a prism; the prism decomposes the ribbon, spreading out its constituent colours like a partially opened fan, and the coloured beam or *spectrum* thus produced is then examined by help of a telescope attached by a movable joint to a stand

which carries the prism and object-tube. Sodium compounds, under these circumstances, give yellow light only, indicated by a double band of light in a position corresponding to a portion of the yellow part of an ordinary solar spectrum. The potassium spectrum is mainly composed of a red and violet band; lithium a crimson, and, at very high temperatures, a blue band. Most of the other elements give equally characteristic spectra.

By aid of a combined microscope and spectroscope (micro-spectroscope) the colour of coloured fluids can be analysed.

CHEMICAL TOXICOLOGY.

In cases of criminal and accidental poisoning, the substances presented to the chemical analyst for examination are usually articles of food, medicines, vomited matters, or the liver, kidney, intestines, stomach and contents, removed in course of post-mortem examination. In these cases some special operations are necessary before the poison can be isolated in a state of sufficient purity for the application of the usual tests; for in most instances the large quantity of animal and vegetable, or, in one word, organic matter present, prevents or masks the characteristic reactions on which the tests are founded. These operations will now be described*; they form the chemical part of the subject of Toxicology (*τοξικὸν, toxicon*, poison, and *λόγος, logos*, discourse).

Substances occurring in the form of an apparently definite salt or unmixed with organic matter need no special treatment (they are analysed by the ordinary methods already given), attention being restricted to poisonous compounds only.

EXAMINATION OF AN ORGANIC MIXTURE SUSPECTED TO CONTAIN:—

MERCURY, ARSENICUM, ANTIMONY, LEAD, COPPER, OR ZINC;
SULPHURIC ACID, NITRIC ACID, HYDROCHLORIC ACID, OXALIC

* Materials for these experiments are readily obtained for educational purposes by dissolving the poison in infusions of tea or coffee, in porter or in water to which some mucilage of starch or linseed meal, pieces of bread, potato, and fat have been added.

ACID, OR HYDROCYANIC ACID; CAUSTIC ALKALIES; PHOSPHORUS; STRYCHNIA, MORPHIA, OR OTHER POISONOUS ALKALOIDS.

Preliminary Examination.

Odour, Appearance, Taste.—Smell the mixture, with the view of ascertaining the presence or absence of any notable quantity of free hydrocyanic acid. Look carefully for any small solid particles, such as arsenic, corrosive sublimate, or verdigris, and for any appearance which may be regarded as abnormal, any character unusual to the coffee, tea, beer, medicine, vomit, coats of stomach, kidney, liver, or other organ, tissue, or solid matter under examination.

Poisonous quantity of acid.—Add to a small portion some solution of carbonate of sodium, with the view of ascertaining by strong effervescence the presence of any large, poisonous quantity of sulphuric, nitric, or hydrochloric acid (p. 581).

Poisonous quantity of alkali.—If so excessively alkaline as to require the addition of a very large quantity of acid before neutralization is effected, a noxious quantity of a corrosive or caustic alkali is present. Whether soda or potash is ascertained by the usual tests.

Special instructions may induce the operator to suspect the presence of one particular poison. Direct examination for the latter may then be made, either at once, if the substance has an aqueous character, or when filtration or treatment with warm hydrochloric or acetic acid has afforded a more or less colourless liquid.

Fluids.—A vomit or the contents of a stomach, if set aside in a long narrow vessel (test-glass or ale-glass), or, better, exposed on a filter during a night, will often yield a more or less clear limpid portion at the bottom or top of the solid matter. This fluid (separated by a pipette or otherwise) will sometimes respond to tests without further preparation, and always requires less preparatory treatment than a semisolid mixture. If none passes through a filter, a portion often collects in the upper part.

General procedure.—If the preliminary examination does

not indicate the method to be pursued, proceed as follows, treating a portion (not more than one-fourth) of the mixture for the poisonous metals, another for the acids, and a third for alkaloids, reserving the remainder for any special experiments which may suggest themselves in the course of analysis.

Examination for Mercury, Arsenicum, Antimony, Lead, Copper, Zinc.

If a liquid, acidulate with hydrochloric acid and boil for a short time. If solid or semisolid, cut up the matter into small pieces, add enough water to form a fluid mixture, stir in ten or twenty per cent. of ordinary liquid hydrochloric acid, and boil until, from partial aggregation and solution of the solid matter, filtration can be easily effected.

Heat a portion of the clear liquid with a thin piece of bright pure copper or copper gauze, about an inch long and a quarter of an inch broad, for about ten or twenty minutes; metallic mercury, arsenicum, or antimony will be deposited on the copper, darkening it considerably in colour. Pour off the liquid from the copper, carefully rinse the latter with a little cold water, dry the piece of metal by holding it over or near a flame (using fingers, not tongs, or it may become sufficiently hot for loss of mercury or arsenicum to occur by volatilization), introduce it into a narrow test-tube or piece of glass tubing closed at one end, and heat the bottom of the tube in a flame, holding it horizontally that the upper part of the tube may be kept cool, and partially closing the mouth with the finger to prevent escape of vapour. Under these circumstances any Mercury will volatilize from the copper and condense on the cool part of the tube in a ring or patch of white sublimate, readily aggregating into visible globules on being pressed by the side of a thin glass rod inserted into the tube; Arsenicum will volatilize from the copper, and, absorbing oxygen from the air in the tube, condense on the cool part of the glass in a ring or patch of white sublimate of arsenic (grey or even darker if much arsenicum as well as arsenic be

present), not running into globules when rubbed, but occurring in small crystals, the characteristic octahedral form of which (*vide* p. 191) is readily seen by aid of a good hand lens, or the low power of a microscope; *Antimony* volatilizes from the copper if strongly heated, and, absorbing oxygen, immediately condenses as a slight white deposit close to the metal.

Confirmatory Tests.—1. Nothing short of the production of globules should be accepted as evidence of the presence of mercury. It will usually have existed as corrosive sublimate.

2. To confirm indications of the presence of arsenicum, a portion of the acid liquid may be subjected to the hydrogen tests (pp. 192 to 195); or the tube containing the white crystalline arsenic may be broken, and the part on which the sublimate occurs boiled for some time in water, and the hydrosulphuric-acid, ammonio-nitrate-of-silver, and ammonio-sulphate-of-copper tests (pp. 196, 197) applied to the aqueous solution.

3. For antimony, a portion of the acid liquid must always be introduced into the hydrogen-apparatus with the usual precautions. (*Vide* p. 207.)

Note.—Before finally concluding that arsenicum is absent from a fluid, the latter should be warmed with a little sulphurous acid or oxalic acid and ordinary tests then again applied; for arsenic acid and other arseniates are not readily affected by the usual reagents for arsenicum.

For *Lead* and *Copper*, pass hydrosulphuric acid gas through the clear acid liquid for some time, warming the liquid if no precipitate is produced, or diluting and partially neutralizing the acid by ammonia if much acid has been added. Collect on a filter any black precipitate that may have formed; wash, dissolve in a few drops of aqua regia, dilute, and apply tests, such as ammonia for copper, sulphuric acid for lead, or any other of the ordinary reagents (pp. 217, 243).

Copper may often be at once detected in a small quantity of acidulated liquid by immersing the point of a penknife or a piece of bright iron wire—a deposit of copper, in its characteristic colour, quickly or slowly appearing, according to the amount present (p. 216).

Zinc.—To the acid liquid through which sulphuretted hydrogen has been passed, add excess of ammonia (or to the original acid fluid add excess of ammonia, and then sulphhydrate of ammonium); a precipitate falls which may contain alumina, phosphates, and zinc—it is usually blackish, from the presence of sulphide of iron. Collect the precipitate on a filter, wash, dissolve in a little hydrochloric acid, add a few drops of nitric acid, boil, pour in excess of ammonia, filter, and test the filtrate with sulphhydrate of ammonium; a white precipitate indicates zinc.

Examination for Mineral Acids, Oxalic Acid, or Hydrocyanic Acid.

To detect *Hydrochloric, Nitric, or Sulphuric Acid* in any liquid containing organic matter, dilute with water and apply to small portions the usual tests for each acid, disregarding indications of small quantities. (*Vide* pp. 310, 334, 362.)

Excessive sourness, copious evolution of carbonic acid gas on the addition of carbonate of sodium, and abundant evidence of acid on applying the various tests to small portions of the fluid presented for analysis, collectively form sufficient evidence of the occurrence of a poisonous amount of either of the three common mineral acids. Small quantities of the hydrochloric, nitric, and sulphuric radicals, occurring as metallic salts or acids, are common normal constituents of food; hence the direction to disregard insignificant indications. If the fluid under examination be a vomit or the contents of a stomach, and an antidote has been administered, free acid will not be found, but, instead, a large amount of the corresponding salt.

For *Oxalic Acid*, filter or strain a portion of the liquid, if not already clear, and add solution of acetate of lead so long as a precipitate occurs; collect the precipitate, which is partly oxalate of lead, on a filter, wash, transfer it to a test-tube or test-glass, add a little water, and pass hydrosulphuric gas through the mixture for a short time; the lead is thus converted into the insoluble form of sulphide, while oxalic acid is set free in the solution. Filter, boil to get rid of hydrosulphuric gas, and apply the usual tests for oxalic acid (*see* p. 369) to the clear filtrate.

The contents of a stomach containing oxalic acid is often of a dark-brown colour with a tinge of green (altered blood and mucus), and the viscid mixture generally, though slowly, affords some clear limpid, almost colourless, liquid by filtration on standing.

For *Hydrocyanic Acid*, the three chief tests may be applied at once to the liquid or semiliquid organic mixture, whether it has an odour of hydrocyanic acid or not. First:—Half fill a small porcelain crucible with the material, add eight or ten drops of strong sulphuric acid, stir gently with a glass rod, and invert over the mouth of the crucible a watch-glass moistened with a small drop of solution of nitrate of silver; a white film on the silver solution is probably cyanide of silver, formed by the action of the gaseous hydrocyanic acid on the nitrate of silver. Second:—Prepare a small quantity of the organic mixture as before, slightly moistening the centre of the watch-glass with solution of potash; here, again, the heat generated by the action of the strong acid is sufficient to volatilize some of the hydrocyanic acid, which, reacting on the potash, forms cyanide of potassium. On removing the watch-glass and stirring into it successively solution of a ferrous salt, a ferric salt, and hydrochloric acid, flocks of prussian blue are produced if hydrocyanic acid is present. Third:—Proceed as before, moistening the watch-glass with sulphurate of ammonium; after exposure to the hydrocyanic gas for five or ten minutes, add a drop of solution of ammonia, evaporate to dryness at a low temperature, and add a drop of hydrochloric acid and of solution of perchloride of iron; a blood-red colour, due to sulphocyanate of iron, is produced if cyanogen is present.

If the above reactions are not well marked, the organic mixture may be carefully and slowly distilled in a small retort, the neck of which passes into a bottle and dips beneath the surface of a little water at the bottom of the bottle, and the reagents then applied to separate portions of the distillate.

The examination of organic mixtures for hydrocyanic acid must be made without delay, as the poison soon begins to decompose, and in a day or two is usually destroyed.

Examination for Phosphorus.

A paste containing phosphorus is commonly employed for destroying vermin. In cases of poisoning the phosphorus is commonly in sufficient quantity to be recognised by its characteristic unpleasant smell. A stomach in which it occurs not unfrequently exhibits slight luminosity if opened in a dark room. When the phosphorus is too small in quantity or too much diffused to afford this appearance, a portion of the material is placed in a flask, water acidulated by sulphuric acid added, a long wide glass tube fitted to the neck of the flask by a cork, and the mixture gently boiled. If phosphorus is present (even 1 part in 2,000,000, according to De Vrij) the top of the column of steam as it condenses in the tube will appear distinctly phosphorescent when viewed in a dark room. From its liability to oxidation phosphorus cannot be detected after much exposure of an organic mixture to air.

Examination for Strychnia and Morphia.

Strychnia.—If solid or semisolid, digest the matter with water and about ten per cent. of hydrochloric acid till fluid, filter, evaporate to dryness over a water-bath. If the organic mixture is already liquid, it is simply acidulated with hydrochloric acid and evaporated to dryness. The acid residue is next treated with spirit of wine as long as anything is dissolved, the filtered tincture evaporated to dryness over the water-bath, and the residue digested in water and filtered. This slightly acid aqueous solution must now be rendered alkaline by ammonia, and well shaken in a closed bottle or long tube with about half an ounce of chloroform, and set by till the chloroform has subsided. The chloroform (which contains the strychnia) is then removed by a pipette, the presence of any aqueous liquid being carefully avoided, and evaporated to dryness in a small basin over a water-bath, the residue moistened with concentrated sulphuric acid, and the basin kept over the water-bath for several hours. (It is highly important that the sulphuric acid used in this operation should be free from nitrous compounds. Test the acid, therefore, by adding

powdered sulphate of iron, which becomes pink if nitrous bodies are present. If these are found, the acid should be purified by strongly heating with sulphate of ammonium, seventy or eighty grains to a pint.) The charred material is exhausted with water, filtered, excess of ammonia added, the filtrate shaken with about a quarter of an ounce of chloroform, the mixture set aside for the chloroform to separate, and the chloroform again removed. If on evaporating a small portion of this chloroform solution to dryness, adding a drop of sulphuric acid to the residue, and warming, any darkening in colour or charring takes place, the strychnia is not sufficiently pure for chemical detection; in that case the rest of the chloroform must be removed by evaporation, and the residue redigested in warm sulphuric acid for two or three hours. Dilution, neutralization of acid by ammonia, and agitation with chloroform is again practised, and the residue of a small portion of the chloroform solution once more tested with sulphuric acid. If charring still occurs, the treatment must be repeated a third time. Finally a part of the chloroform solution is taken up by a pipette and drop after drop evaporated on one spot of a porcelain crucible-lid until a fairly distinct dry residue is obtained. A drop of sulphuric acid is placed on the spot, another drop placed near, a minute fragment of red chromate of potassium placed in the second drop, and, when the acid has become tinged with the chromate, one drop drawn across the other; the characteristic evanescent purple colour is then seen, if strychnia is present. Other tests (*vide* p. 460) may be applied to similar spots.

This is Girdwood and Rogers's method for the detection of strychnia when mixed with organic matter. It is tedious but trustworthy, and, though apparently complicated, very simple in principle, thus;—strychnia is soluble in acidulated water or alcohol, or in chloroform, readily removed from an alkaline liquid by agitation with chloroform, and not charred or otherwise attacked when heated to 212° F. with sulphuric acid: much of the organic matter of the food is insoluble in water; of that soluble in water, much is insoluble in alcohol; and of that soluble in both menstrua, all is charred and destroyed by warm sulphuric acid in a shorter or longer time.

Morphia and the Meconic Acid with which it is associated in Opium.—To the liquid or the semifluid mixture warmed for some time with a small quantity of acetic acid, filtered, and concentrated if necessary, add solution of acetate of lead until no further precipitate is produced. Filter and examine the precipitate for meconic acid, reserving the filtrate for the detection of morphia.

The Precipitate.—Wash the precipitate (meconate of lead, etc.) with water, place it in a test-tube or test-glass with a small quantity of water, pass hydrosulphuric acid gas through the mixture for a short time, filter, slightly warm in the small basin, well stirring to promote removal of excess of the gas, and add a drop of neutral solution of perchloride of iron; a red colour, due to the formation of meconate of iron, is produced if meconic acid is present. This colour is not destroyed on boiling the liquid, as is the case with ferric acetate, nor is it bleached by solution of corrosive sublimate, thus distinguishing it from ferric sulphocyanate. It is discharged by hydrochloric acid.

The Filtrate.—The solution from which meconic acid has been removed by acetate of lead is evaporated to a small bulk over a water-bath, excess of carbonate of potassium added, and evaporation continued to dryness. The residue is then treated with alcohol, which dissolves the morphia. The alcoholic solution evaporated similarly may leave the morphia sufficiently pure for the application of the usual tests (*vide* p. 450) to small portions of the residue. If no reaction is obtained, add a drop of sulphuric acid and a little water to the residue and shake with ether, in which the salt of morphia is insoluble. The treatment with ether may be repeated until nothing more is removed, the acid aqueous liquid saturated with carbonate of potassium, the mixture evaporated to dryness, the residue digested in alcohol, filtered, and portions of the alcoholic liquid evaporated to obtain spots of morphia for the application of the ordinary tests.

If much organic matter is believed to remain in the filtrate after the acetate of lead treatment, or if a considerable excess

of acetate of lead has been employed, the filtered liquid should be subjected to a current of sulphuretted hydrogen until no more sulphide of lead is precipitated, the mixture filtered, and the filtrate, with the washings from the sulphide of lead, evaporated to a small bulk, excess of carbonate of potassium added, the whole well mixed and agitated with twice or thrice its bulk of a mixture of ether and acetic ether (ether alone might not dissolve the morphia). On standing, the ethereal liquid rises to the surface: it is carefully removed, evaporated to dryness, and the residue tested or further purified in the manner described in the preceding paragraph.

The examination for morphia must be conducted with great care, and with as large a quantity of material as can be spared; for its isolation from other organic matter is an operation of considerable difficulty, especially when only a minute proportion of alkaloid is present. Fortunately the detection of meconic acid does not include similar difficulties; and as its reactions are quite characteristic, its presence is held to be strong evidence of the existence of opium in an organic mixture.

Examination for other Poisonous Alkaloids.

Stas's Process.—Minutely subdivide any solid matter; to this and the liquid portion of the vomit, etc., add about twice their weight of the strongest spirit of wine containing sufficient tartaric acid to fairly acidify the mixture. Digest the whole in a flask at a temperature of 150° or 160° F.; set aside to cool; filter. The solution, which will contain the whole of the alkaloid, should then be evaporated nearly to dryness *in vacuo*, or at all events at a temperature not exceeding 100° F., lest volatile alkaloids should be dissipated. The residue is next exhausted with cold anhydrous alcohol; filtered; and the filtrate evaporated to dryness with the precautions already stated. The extract is dissolved in a very small quantity of water, treated with excess of powdered bicarbonate of sodium or potassium, and well shaken with five or six times its volume of pure ether (with perhaps a little acetic ether). This ethereal liquid contains the alkaloid. Small portions should be evaporated in watch-glasses and

tasted, or tested physically and chemically, according as the knowledge of collateral circumstances by the operator, or his experience, or such reactions as are recorded on pp. 462-472, may suggest.

If a volatile alkaloid (conia, nicotia, hyoscyamia, lobelina) is indicated, the ethereal solution, which may still contain animal matter, is removed, agitated with aqueous solution of potash, decanted, and shaken with pure diluted sulphuric acid. On standing, the aqueous portion, containing the alkaloid as acid sulphate, subsides; the upper ethereal portion containing the animal matter is rejected; the acid aqueous liquid is made alkaline with caustic potash or soda; ether added; well shaken; the ethereal liquid decanted, evaporated to dryness *in vacuo*, or at a low temperature, and (to get rid of all traces of ammonia) again moistened with ether and dried. The residue is now tested for the suspected alkaloid by taste, smell, and the application of appropriate reagents (pp. 462-472).

If a non-volatile alkaloid (aconitia, atropia, brucia, colchicia, emetia, physostigmia, solania, veratria, as well as morphia, codeia, and strychnia) is indicated, further purification is effected by decanting the ethereal liquid from the lower aqueous solution of bicarbonate of sodium, removing the ether by evaporation, digesting the residue in alcohol, filtering, evaporating the alcohol, treating the residue with dilute sulphuric acid, setting aside for a few hours, filtering, concentrating, adding powdered carbonate of potassium, and finally anhydrous alcohol. The alcoholic liquid, on evaporation, yields the alkaloid in a fit state for testing in the manner already stated.

Sonnenschein's Process.—Digest with diluted hydrochloric acid, evaporate to the consistence of syrup, dilute, set aside for some hours, filter. Add solution of phosphomolybdic acid so long as any precipitate falls or cloudiness occurs; collect the precipitate on a small filter; wash it with water containing phosphomolybdic and nitric acid, and, while still moist, place it in a flask. Decompose this compound of phospho-

molybdic acid and alkaloid by adding caustic baryta until the stirred mixture is distinctly alkaline. Distil off volatile alkaloids, condensing and collecting by help of a long tube, so bent that the apparatus shall act as a retort, the end of the tube being attached to a bulb or a series of bulbs containing dilute hydrochloric acid. The acid liquid evaporated gives a residue of hydrochlorates of alkaloids. The latter will afford characteristic reactions with the tests for the suspected alkaloid, and on being moistened with baryta-water and warmed, will afford fumes of volatile alkaloids whose odour is usually characteristic. The residue in the flask will contain non-volatile alkaloids. It is treated with carbonic acid gas to neutralize and precipitate the excess of baryta as insoluble carbonate of barium; the mixture is evaporated to dryness over a water-bath; and the residue digested in alcohol. The alcoholic solution evaporated generally yields the alkaloids in a fit state for testing.

Reagents for Alkaloids.

Phosphomolybdic acid forms with ammonia, in acid solutions, a remarkably insoluble compound, and it comports itself in a similar manner with those compounds which are analogous to ammonia—the nitrogenized organic bases—consequently forming an excellent reagent for their detection. It may be prepared in the following manner:—Molybdate of ammonium is precipitated by phosphate of sodium; the yellow precipitate, having been washed, is diffused through water, and heated with sufficient carbonate of sodium to dissolve it. The solution is then evaporated to dryness, and calcined to drive off the ammonia. In case any of the molybdic compound be reduced by this operation, the residue must be moistened with nitric acid and again calcined. The dry mass is then dissolved in cold water, the solution strongly acidulated with nitric acid, and water added until ten parts of the solution contain one of the dry salt. The liquid, which is of a golden-yellow colour, must be preserved from ammoniacal fumes. It precipitates all the alkaloids (with the exception of urea) when a mere trace only is present. The precipitates are yellow, generally flocculent, insoluble in water, alcohol, ether, and the dilute mineral acids, with the

exception of phosphoric acid. Nitric, acetic, and oxalic acids, concentrated and boiling, dissolve them. These compounds are decomposed by the alkalies, certain metallic oxides, and the alkaline salts, which separate the alkaloid. To give an idea of the sensitiveness of this reagent, it may be stated that the 0.000071 gramme of strychnia gives an appreciable precipitate with one cubic centimètre of the solution of phosphomolybdic acid.

Other special reagents for alkaloids are the "Nessler test" (see Index) and a solution of the double "Iodide of Bismuth and Potassium." The latter is made (by Thresh) on adding together one ounce of *Liquor Bismuthi*, B. P., 90 grains of iodide of potassium, and 90 grains of strong hydrochloric acid. This orange-coloured solution gives a red precipitate with dilute cold solutions containing alkaloids.

Obscure Poisons.—Many substances, the active principles of which are at present beyond the reach of the chemical analyst, are poisons of a more or less active character. (See the *Pharmaceutical Journal* for Sept. 6, 1879, p. 195, and for Dec. 20, 1879, p. 481.)

ANTIDOTES.

Vide "Antidotes" in the Index.

QUESTIONS AND EXERCISES.

874. In examining food and similar matter for poison, why must not the ordinary tests for the poison be at once applied?

875. What preliminary operations should be performed on a vomit in a case of suspected poisoning?

876. How would you proceed in searching for corrosive sublimate in wine?

877. By what series of operations would you satisfy yourself of the presence or absence of arsenic in the contents of a stomach?

878. Describe the treatment to which decoction of coffee should be subjected in testing it for tartar-emetic.

879. State the method by which the occurrence of lead in water is demonstrated.

880. Give a process for the detection of copper in jam.

881. How would you detect zinc in a vomit?

882. How may the presence of a poisonous quantity of sulphuric acid in gin be proved ?

883. In examining ale for free nitric acid what reactions would be selected ?

884. Show how you would conclude that a dangerous quantity of hydrochloric acid had been added to cider.

885. Describe the manipulations necessary in testing for hydrocyanic acid in the contents of a stomach.

886. By what method is oxalic acid discovered in infusion of coffee ?

887. How is phosphorus detected in organic mixtures ?

888. Give the process by which strychnia is isolated from partially digested food.

889. Mention the experiments by which the presence of laudanum in porter is demonstrated.

890. Name the appropriate antidotes in cases of poisoning by : —*a*, alkaloids ; *b*, antimonials ; *c*, arsenic ; *d*, barium salts ; *e*, copper compounds ; *f*, hydrochloric acid ; *g*, hydrocyanic acid ; *h*, preparations of lead ; *i*, corrosive sublimate ; *j*, nitric acid ; *k*, oxalic acid ; *l*, salts of silver ; *m*, oil of vitriol ; *n*, tin liquors ; *o*, zinc solutions ; *p*, carbolic acid.

EXAMINATION OF MORBID URINE AND CALCULI.

The various products of the natural and continuous decay of animal tissue and the refuse matter of food are eliminated from the system chiefly as fæces, urine, and expired air. Air exhaled from the lungs carries off from the blood much carbon (about 8 ounces in 24 hours), in the form of carbonic acid gas, and some aqueous vapour—the latter, together with a small amount of oily matter, also escaping by the skin. Directing the breath to a cold surface renders moisture evident ; and breathing through a tube into lime-water demonstrates the presence of a considerable quantity of carbonic acid gas. The fæces consist mainly of the insoluble *débris* of the system, the soluble matters and water forming the urine. These excretions vary considerably, according to the food and general habits of the individual and external temperature. But in disease the variations become excessive ; their detec-

tion by the medical practitioner, or by the pharmacist for the medical practitioner, is therefore a matter of importance.

A complete analysis of fæces, urine, or expired air cannot be performed in the present state of our knowledge. Nor can even a partial analysis of fæces or air be made with sufficient ease and rapidity to be practically available in medical diagnosis. But with regard to urine, certain abnormal substances and abnormal quantities of normal constituents may be chemically detected in the course of a few minutes by any one having already some knowledge of chemical manipulation.

Healthy human urine contains, in 1000 parts, 957 of water, 14 of urea, 1 of uric acid, 15 of other organic matter, and 13 of inorganic salts. The amount passed in twenty-four hours varies from two to three pints in an adult, and its specific gravity, if healthy, will range from 1.015 to 1.025. The acidity of urine Thudichum considers to be due to *cryptophanic acid*, $\text{H}_2\text{C}_5\text{H}_7\text{NO}_5$.

EXAMINATION OF MORBID URINE FOR ALBUMEN, SUGAR, BILE,
AND EXCESS OF UREA; AND URINARY SEDIMENT FOR URATES
(OR LITHATES), PHOSPHATES, OXALATE OF CALCIUM, AND
URIC ACID.

Albumen.—To detect albumen, acidulate a portion of the clear urine in a test-tube with a few drops of acid (to keep phosphates in solution: nitric is best, acetic not so good) and boil; flocks or coagula will separate if albumen be present.

This experiment should be first made on normal urine containing a drop or two of solution of white of egg. The coagulum is white if it is only albumen, greenish if bile-pigment be present, and brownish-red if the urine contain blood. The influence of acids and alkalies on the precipitation of albumen is noticed on page 536.

The occurrence of albumen in the urine may be temporary and of but little importance; or it may indicate the existence of a serious affection, known as Bright's disease.

Sugar.—To a portion of the clear urine in a test-tube add five or ten drops of solution of sulphate of copper; pour in solution of potash or soda until the precipitate first formed is redissolved; slowly heat the solution to near the boiling-

point; a yellow, yellowish-red, or red precipitate (cuprous oxide) is formed if sugar be present.

This experiment should be first made on urine containing a drop or two of solution of grape-sugar (p. 484). The hydrate of copper precipitated by the alkali is insoluble in excess of pure potash or soda, but readily dissolves if organic matter, especially sugar, be present. The copper salt should not contain iron.

Other tests may be applied if necessary (*vide* p. 487).

A minute amount of sugar is said to occur in normal urine, and a distinct trace is occasionally present. In larger quantities (often 5 per cent.) it is a characteristic constituent of the urine of diabetic patients, greatly increasing the specific gravity of the excretion. Small hydrometers (termed urinometers) are commonly employed for quickly and readily ascertaining the specific gravity of urine; they range from 1.000 to 1.050, the interval of 1.015 to 1.025 being marked as "H. S." or "healthy state." (*Vide* "Specific Gravity" and "Hydrometers" in Index.)

Bile.—This is best detected by the general test (Pettenkofer's) described on page 545. Or a little of the urine may be placed on a white plate and strong nitric acid dropped on it; a peculiar play of colours—green, yellow, violet, etc.—occurs if (the colouring-matter of) bile be present.

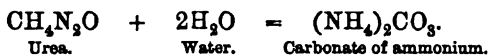
Excess of Urea.—About one-third of the solid matter in the urine is urea. Its proportion varies considerably; but $1\frac{1}{2}$ per cent. may be regarded as an average amount. Concentrate urine slightly by evaporation in a small dish, pour the liquid into a test-tube, set the tube aside till cold, or cool it by letting cold water run over the outside, add an equal bulk of strong nitric acid and again set aside; scaly crystals of nitrate of urea are deposited more or less quickly.

With regard to the amount of urea in urine, it is impossible to sharply define excess or deficiency. If nitric acid gives crystals without concentration, excess is certainly present. A rough estimate may be formed by mixing a few drops of the urine and acid on a piece of glass and setting aside; the time which elapses before crystals form is an indication of the quantity in the specimen. The time will vary according to the temperature and state of moisture of the atmosphere;

but with care some useful comparative results may in this way be obtained.

(For methods of estimating the amount of urea in urine, and for drawings of the necessary apparatus, *vide* "Journal of the Chemical Society," 1874, p. 749; and 1877, pp. 534 and 538; also *Chemical News*, Jan. 22, 1875.)

Tests.—Urea in solution in water may be detected by the above reaction with nitric acid, and by the readiness with which it yields ammonia on being boiled with alkalis. In putrid urine its conversion into an ammoniacal salt has already been effected by ammoniacal fermentation.



This transformation of the urea into carbonate of ammonium is due to the action of a special ferment belonging to the genus *Torulacei*, formed of chaplets of globules similar in form but much smaller than those of beer yeast. It occurs as a white deposit in the urine. If some of this deposit be added to a saccharine solution containing urea it rapidly multiplies, carbonate of ammonium being formed.

Formula of Urea.—The empirical formula of urea is $\text{CH}_4\text{N}_2\text{O}$.

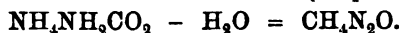
Its rational formula may be thus written:— $\left. \begin{array}{c} (\text{CO})'' \\ \text{H}_2 \\ \text{H}_2 \end{array} \right\} \text{N}_2$; that is,

it may be regarded as one of the organic bases already referred to, a primary diamine, in which the bivalent radical CO occupies the place of H_2 . The other atoms of hydrogen may be displaced by various radicals, and many *compound ureas* thus be obtained.

Artificial Urea.—Urea may be prepared artificially by Williams's modification of Wöhler's method. Cyanide of potassium, of the best commercial quality (containing about 90 per cent. of real cyanide), is fused at a very low red heat in a shallow iron vessel; red lead is added in small quantities at a time, the temperature being kept down by constant stirring. When the red lead ceases to cause further action the mixture (cyanate of potassium and lead) is allowed to cool, the product finely powdered, exhausted with cold water, nitrate of barium added till no more precipitate (carbonate of barium) falls, the mixture filtered, and the filtrate treated with nitrate of lead so long as cyanate of lead is thrown down. The

latter is thoroughly washed, and dried at a low temperature. Equivalent quantities of cyanate of lead and sulphate of ammonium digested in a small quantity of water at a gentle heat (*vide* p. 395) and filtered yield a solution from which urea crystallizes on cooling.

Another process.—Basaroff has found that urea is produced when ordinary carbonate of ammonium is heated in hermetically sealed tubes to about 275° F. for a few hours. The same chemist had previously obtained urea by similarly heating pure carbamate of ammonium; so that the source of the urea in the former case is probably the carbamate of ammonium believed to occur in the carbonate (*see* p. 98).



URINARY SEDIMENTS.

Warm the sediment with the supernatant urine and filter.

<i>Insoluble.</i> Phosphates, oxalate of calcium, and uric acid. Warm with acetic acid, and filter.		<i>Soluble.</i> Urates—of ammonium, calcium, or sodium, chiefly the latter. They are redeposited as the liquid cools, and if sufficient in quantity may be further examined for ammonium, calcium, sodium, and the uric radical by the appropriate tests.
<i>Insoluble.</i> Oxalate of calcium and uric acid. Warm with hydrochloric acid, filter.	<i>Soluble.</i> Phosphates. Add ammonia, white ppt. = phosphate of calcium, or ammonio-magnesium phosphate, or both.	
<i>Insoluble.</i> Uric acid. Apply murexid test (p. 422).	<i>Soluble.</i> Oxalate of calcium. May be reprecipitated by ammonia.	

Notes.—Urinary deposits are seldom of a complex character: the action of heat and acetic and hydrochloric acids generally at once indicates the character of the deposit, rendering filtration and precipitation unnecessary.

The urates are often of a pink or red colour, owing to the

presence of a pigment termed *purpurine*; hence the common name of *red gravel* for such deposits. *Purpurine* is soluble in alcohol, and may be removed by digesting a red deposit in that solvent. It is seldom necessary to determine whether the urate be that of ammonium, calcium, or sodium (*see also Uric Acid*, page 422).

The *phosphate of calcium* and the *ammonio-magnesium phosphate* are usually both present in a phosphatic deposit, the magnesium salt forming the larger proportion. They may, if necessary, and if sufficient in quantity, be separated by collecting on a filter, washing, and boiling with solution of carbonate of sodium. The carbonates of calcium and magnesium thus formed are collected on a filter, washed, and dissolved in a drop or two of hydrochloric acid; chloride of ammonium, ammonia, and carbonate of ammonium are added, and the mixture boiled and filtered; any calcium originally present will then remain insoluble as carbonate of calcium; while any magnesium will be precipitated from the filtrate as ammonio-magnesium phosphate on the addition of phosphate of sodium, the mixture being also well stirred.—The chief portion of excreted phosphates is carried off by the *fæces*, that remaining in the urine being kept in solution by the influence of acid phosphate of sodium and, frequently, lactic acid.—Occasionally, an hour or two after a hearty meal, the urine becomes sufficiently alkaline for the phosphates to be deposited, and the urine when passed is turbid from their presence.—The ammoniacal constituent of the magnesium salt does not occur normally, but is produced from urea as soon as urine becomes alkaline.

Oxalate of calcium is seldom met with in excessive amounts, but very often in small quantities mixed with phosphates. In one case of oxaluria the whole urine excreted by a patient in twenty-four hours furnished to the author only two-thirds of a grain of oxalate of calcium.

Free uric acid is in most cases distinctly crystalline, and nearly always of a yellow, red, or brown colour.

Artificial sediments.—For educational practice, artificial deposits may be obtained as follows:—1. Rub up in a mortar a few grains of serpent's excrement (chiefly urate of ammonium) with an ounce or two of urine; this represents a sediment of urates. 2. Add a few drops of solution of chloride of calcium and of phosphate of sodium to urine; the deposit may be regarded as one of phosphates. 3. To an

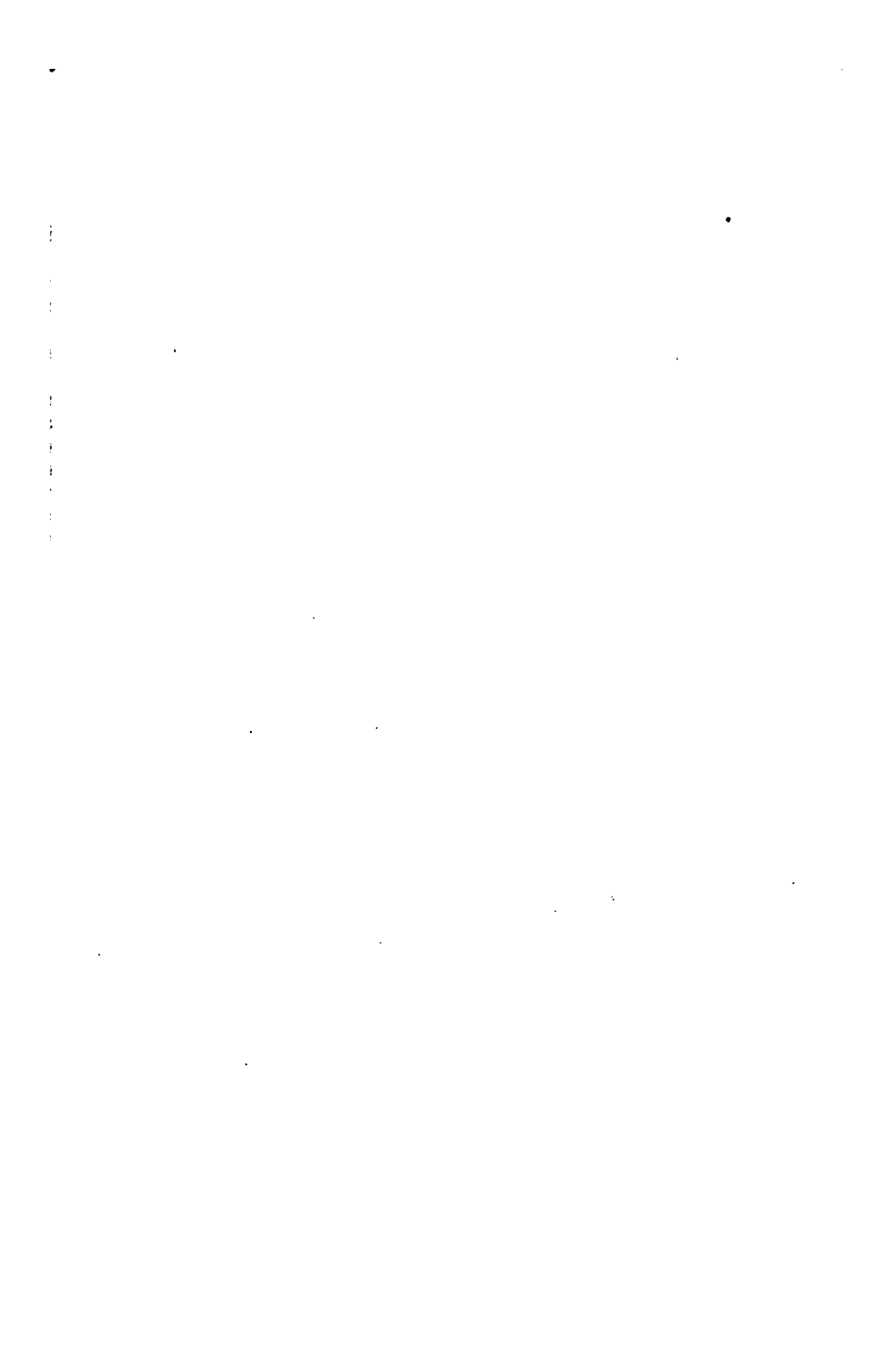
ounce or two of urine add very small quantities of chloride of calcium and oxalate of ammonium; the precipitate is oxalate of calcium. 4. To urine acidulated by hydrochloric acid add a little serpent's excrement; the sediment is uric acid.

Other deposits than the foregoing are occasionally observed. Thus *hippuric acid* ($\text{HC}_9\text{H}_8\text{NO}_3$), a normal constituent of human urine, and largely contained in the urine of herbivorous animals, is sometimes found associated with uric acid in urinary sediment, especially in that of patients whose medicine contains benzoic acid (p. 396). Its appearance, as observed by aid of the microscope, is characteristic—namely, slender, four-sided prisms, having pointed ends. *Cystin* ($\text{C}_3\text{H}_7\text{NSO}_2$) (from *cystis*, *küstie*, a bladder, in allusion to its origin) rarely occurs as a deposit in urine. It is not soluble in warm urine or dilute acetic acid, and scarcely in dilute hydrochloric acid—hence would be met with in testing for free uric acid. It is very soluble in ammonia, recrystallizing from a drop of the solution placed on a piece of glass in characteristic microscopic six-sided plates. *Organized sediments* may be due to the corpuscles of pus, mucus, or blood, fat-globules, spermatozoa, cylindrical casts of the tubes of the kidneys, epithelial cells from the walls of the bladder, or foreign matters, such as fibres of wool, cotton, small feathers, dust, starch; these are best recognised by the microscope. (See accompanying Plates, and the following paragraphs on the microscopic appearances of both crystalline and organized urinary sediments.)

MICROSCOPIC EXAMINATION OF URINARY SEDIMENTS.

Urine containing insoluble matter is usually more or less opaque. For microscopical examination a few ounces should be set aside in a conical test-glass for an hour or two, the clear supernatant urine poured off from the sediment as far as possible, a small drop of the residue placed on a slip of glass and covered with a piece of thin glass, and examined under the microscope with different magnifying-powers.

The respective appearances of the various crystalline and organized matters are given in the annexed Plates (figs. 50 to 61), the figures of which were kindly drawn by H. B. Brady, F.R.S., from natural specimens in the collections of St. Bartholomew's Hospital, Dr. Sedgwick,



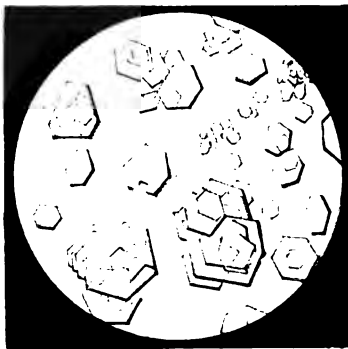
PL
URINARY DEPOSITS (UNORGANIZED)



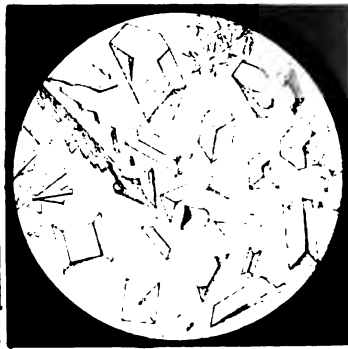
URIC ACID



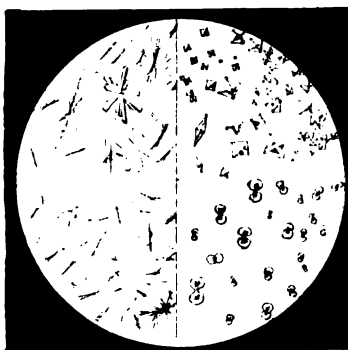
URIC ACID



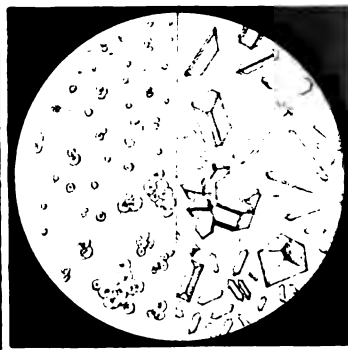
CYSTIN



TRIPLE PHOSPHATE



URATES, a. of Sodium, b. of Magnesium. OXALATE of CALCIUM. CARB. OF CALCIUM. HIPPURIC ACID



As seen with a Zeiss objective & N°1 eye piece (10x) (field of view diam.)

W. W. Stoddart, F.C.S., Mr. Waddington, and the Author.

Uric acid occurs in many forms, most of which are given in the first Plate. Flat, more or less oval crystals, sometimes attached to each other, their outline then resembling an 8, a cross or a star, are common. Single and grouped quadratic prisms, aigrettes, spicula, and crystals recalling dumbbells are met with. From urine acidulated by hydrochloric acid, square crystals, two opposite sides smooth and two jagged, are generally deposited: acidulated by acetic acid more typical forms are obtained. A drop of solution of potash or soda placed on the glass slip will dissolve a deposit of uric acid, a drop of any acid reprecipitating it in minute but characteristic crystals.

Cystin is very rarely met with as a urinary deposit; that from which the figure on Plate I. was taken was found in the urine of a patient in St. Bartholomew's Hospital. Lamellæ of cystin always assume an hexagonal character; but the angles are sometimes ill defined and the plate superposed: in the latter case, a drop of solution of ammonia placed on the glass at once dissolves the deposit, well-marked six-sided crystals appearing as the drop dries up.

Triple Phosphate (phosphate of magnesium and ammonium) is deposited as soon as urine becomes alkaline, the ammoniacal constituent being furnished by the decomposition of urea. It occurs in large prismatic crystals, forming a beautiful object when viewed by polarized light,—sometimes also in ragged stellate or arborescent crystals, resembling those of snow. Both forms may be artificially prepared by adding a small lump of carbonate of ammonium to a few ounces of urine and setting aside in a test-glass.

Amorphous deposits are either *earthy phosphates* (a mixture of phosphates of magnesium and calcium) or *urates* of calcium, magnesium, ammonium, potassium, or sodium—chiefly the latter. They may be distinguished by the action of a drop of acetic acid placed near the sediment on the glass slip, the effect being watched under the microscope; phosphates dissolve, while urates gradually assume characteristic forms of uric acid. Urates redissolve when warmed with the supernatant urines.

Urates of Sodium and Magnesium, though generally amorphous, occasionally take a crystalline form—bundles or tufts of small needles—as shown in the Plate.

Oxalate of Calcium commonly occurs in octahedra requiring high magnifying-power for their detection. The crystals are easily overlooked if other matters are present, but are more distinctly seen after phosphates have been removed by acetic acid. In certain aspects the smaller crystals look like square plates traversed by a cross. A dumbbell form of this deposit is also sometimes seen, resembling certain forms of uric acid and the coalescing spherules of a much rarer sediment—carbonate of calcium. Oxalate of calcium is insoluble in acetic but soluble in hydrochloric acid. The octahedra are frequently met with in the urine of persons who have partaken of garden rhubarb. The crystals may often be deposited artificially (according to Waddington) by dropping a fragment of oxalic acid into several ounces of urine and setting aside for a few hours.

Carbonate of Calcium is rarely found in the urine of man, but frequently in that of the horse and other herbivorous animals. Human urine containing carbonate of calcium often reddens litmus-paper; and it is only after the removal, on standing, of the excess of carbonic acid that the salt is deposited. It consists of minute spherules, varying in size, the smaller ones often in process of coalescence. The dumbbell-form thus produced is easily distinguished from similar groups of uric acid or oxalate of calcium by showing a black cross in each spherule when viewed by polarized light. Acetic acid dissolves carbonate of calcium, liberating carbonic acid gas, with visible effervescence (under the microscope) if the slide has been previously warmed and a group of crystals be attacked.

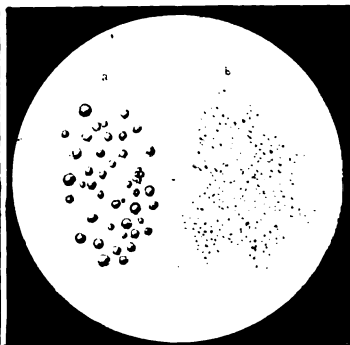
Hippuric Acid.—The pointed rhombic prisms and acicular crystals are characteristic, and easily recognised. The broader crystals may possibly be mistaken for triple phosphate, and the narrower for certain forms of uric acid; but insolubility in acetic acid distinguishes them from the former, and solubility in alcohol from the latter. These tests may be applied while the deposit is under microscopic observation. An alcoholic solution of hippuric acid evaporated to dryness, and the residue treated with water, gives a solution from which characteristic crystalline forms of hippuric acid may be obtained on allowing a drop to dry up on a slip of glass.

Plate II. The organized deposits in urine entail greater care in their determination, and usually require a higher magnifying-power for their proper examination, than

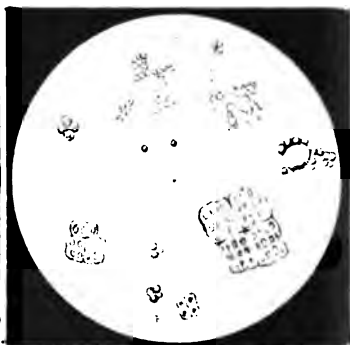
73 IL



BLOOD PROPERTIES



FAT MOLECULES.



SARINA VENTRUCI

Why did you not tell me this?

those of crystalline form. The figures on the second of the accompanying Plates are drawn to 230 diameters. The following notes will assist the observer.

Casts of uriniferous tubuli are fibrinous masses of various forms, and often of considerable length—sometimes delicate and transparent, occasionally granular, and often beset with fat-globules. Epithelial *débris* are frequently present in urine in the form of nucleated cells, regular and oval when full, but angular and unsymmetrical when partially emptied of their contents—sometimes perfect, but more frequently a good deal broken up.

Blood is easily recognised. Urine containing it is high-coloured, and the corpuscles appear under the microscope as reddish circular disks, either single or laid together in strings resembling piles of coin. Their colour and somewhat smaller size serve to distinguish them from pus-corpuscles. In doubtful cases a minute drop of blood taken from a finger by help of a needle should be diluted with water and used for comparison. After urine containing blood has stood for some time, the corpuscles lose their regular rounded outline and become angular. (*See a in the figure.*)

Dr. Day, of Geelong, tests for blood in urine or in stains on clothing by adding a few drops of recently prepared alcoholic solution of the inner unoxidized portions of guaiacum resin, and then a small quantity of Robbins's aqueous or ethereal solution of peroxide of hydrogen, when a blue colour results. "If the stain is on a dark-coloured fabric, the moistened parts may be pressed with white blotting-paper, when blue impressions will be obtained. Contact with many substances causes the blue reaction or oxidation of guaiacum: the peculiarity of blood is that it does not produce this effect unless peroxide of hydrogen or a similar 'antozonic' liquid is present. Bodies such as permanganate of potassium, whose oxygen is apparently in the form of ozone, also cause the production of a blue colour with guaiacum; peroxide of hydrogen and other compounds whose oxygen is in the opposite, positive, or, according to Schönbein, antozonic condition, produce no such effect. It would seem as if blood or some constituent of blood has the power of converting positive into negative oxygen, and thus brings about an effect which negative oxygen alone is able to produce; for of all substances which, like blood, do not alone cause guaiacum to become blue, blood is the only one that so affects 'antozonides' (themselves inactive) as to

enable them to act as ozonides, that is to oxidize the guaiacum. Both the venous and arterial fluid from any red-blooded animal will produce this blue reaction. Fruit-stains are darkened by ammonia, which does not alter the colour of blood. Iron-stains or iron-moulds yield no colour to water, whereas the red colouring-matter of blood is soluble in water. The peroxide of hydrogen should be free from more than a trace of acid."

The blood-corpuscles of ordinary animals are much smaller than those of man; but a $\frac{1}{6}$ - or $\frac{1}{8}$ -inch lens is necessary for proper differentiation (J. G. Richardson).

Pus and Mucus.—Purulent urine deposits, on standing, a light-coloured layer, easily diffused through the liquid by shaking. Acetic acid does not dissolve the sediment; and solution of potash of official strength converts it into a gelatinous mass. Under the microscope, pus-corpuscles appear rounded and colourless, rather larger than blood-disks, and somewhat granular on the surface. They generally show minute nuclei, which are more distinctly seen after treatment with acetic acid. (See the portion of the figure marked a.) Mucus possesses no definite microscopic characters, but commonly has imbedded in it pus, epithelium, and air-bubbles. Mucus is coagulated in a peculiar and characteristic manner by acetic acid; and this reaction, together with the ropy appearance it imparts to urine, prevents it being confounded with pus.

Day's test for pus consists in adding a drop or two of oxidized tincture of guaiacum to the urine or other liquid, when a clear blue colour is produced. It is necessary to moisten dry pus with water before applying the test. The test-liquid is made by exposing a saturated alcoholic solution of guaiacum to the air until it has absorbed a sufficient quantity of oxygen to give it the property of turning green when placed in contact with iodide of potassium. Day's test for mucus consists in the application, first, of oxidized tincture of guaiacum, which by itself undergoes no change in the presence of mucus, and then in the addition of carbolic acid or creasote, which quickly changes the colour of the guaiacum to a bright blue. Neither carbolic acid or creasote alone will render guaiacum blue. In testing for mucus on cloths, or when it is mixed with blood, it is necessary to use the carbolic acid pure; but when the mucus is in a liquid state, it is better to use carbolic acid diluted with alcohol.

Saliva.—This secretion is an aqueous fluid containing less than 1 per cent. of solid matter, of which one-third is an albumenoid substance termed *ptyalin* (from πτύαλον, *spittle*), a body that has the power of converting starch into dextrin and grape-sugar: alkaline salts, including a trace of sulphocyanide of potassium and calcareous compounds, are also present.

Day's test for saliva in urine, etc., is similar to that for mucus, with the exception that the blue reaction produced by the oxidized tincture of guaiacum and alcoholic solution of carbolic acid is highly intensified by the addition of Robbins's ethereal or aqueous solution of peroxide of hydrogen.

Fatty matter occurs either as minute globules partially diffused through the urine (as shown at *a*) or in more intimate emulsion (as at *b* in the figure). When present in larger quantity, it collects as a sort of skim on the surface after standing.

Spermatozoa are liable to escape notice, on account of their small size and extreme transparency. Suspected urines should be allowed to settle some hours in a conical test-glass, and the drop at the bottom examined under a high power. The drawing shows their tadpole-like appearance.

Sarcina ventriculi is an alga of very rare occurrence in urine, though not unfrequent in the matters vomited during certain diseases of the stomach. The upper figures (*a*) are copied from Dr. Thudichum's drawing (from urine); the larger fronds (*b*) are from vomited matter.

Extraneous bodies, such as hair, wool, or fragments of feathers, are often found in urinary deposits; and ludicrous mistakes have been made by observers not on their guard in respect to such casual admixtures.

EXAMINATION OF URINARY CALCULI.

The term *calculus* is the diminutive of *calx*, a lime- or chalk-stone.

Knowledge of the composition of a calculus or urinary deposit affords valuable diagnostic aid to the physician; hence the importance of a correct analysis of these substances.

Nature of Calculi.—Urinary calculi have the same composition as unorganized urinary sediments. They consist, in short, of sediments that have been deposited slowly within the bladder, particle on particle, layer on layer, the several

substances becoming so compact as to be less easily acted on by reagents than when deposited after the urine has been passed—the urates less readily soluble in warm water, the calcic phosphate insoluble in acetic acid until it has been dissolved in hydrochloric acid and reprecipitated by an alkali.

Preliminary treatment.—If the calculus is whole, saw it in two through the centre, and notice whether it is built up of distinct layers or apparently consists of one substance. If the latter, use about a grain of the sawdust for analysis; if the former, carefully scrape off portions of each layer, and examine them separately. If the calculus is in fragments, select fair specimens of about half a grain or a grain each, and reduce to a fine powder by placing on a hard surface and crushing under the blade of a knife.

Analysis.—Commence the analysis by heating a portion, about the size of a pin's head, on platinum foil in order to ascertain whether organic matter, inorganic matter, or both, are present. If both, the ash is examined for inorganic substances, and a fresh portion of the calculus for uric acid by the murexid test. (In the absence of uric acid any slight charring may be considered to be due to indefinite animal matter.) If composed of organic matter only, the calculus will in nearly all cases be uric acid, the indication being confirmed by applying the murexid test in a watch-glass to another fragment, half the size of a small pin's head. If inorganic only, the ash on the platinum foil may be examined for phosphates, and a separate portion of the calculus for oxalates. Even a single drop of liquid obtained in any of these experiments may be filtered by placing it on a filter not larger than a sixpence and previously moistened with water, and adding three or four drops of water one after the other as each passes through the paper. If the calculus is suspected to contain more than one substance, boil about half a grain of the powder in half a test-tubeful of distilled water for a few minutes and pour it on a small filter; then proceed according to the following Table:—

<i>Insoluble.</i> Phosphates, oxalate of calcium, and free uric acid. Boil with two or three drops of hydrochloric acid and filter.		<i>Soluble.</i> Urates. These will probably be redeposited as the solution cools. Small quantities may be detected by evaporating the solution to dryness. They are tested for ammonium, sodium, calcium, and the uric radical by the appropriate reagents.
<i>Insoluble.</i> Uric acid. Apply the murexid test (p. 422).	<i>Soluble.</i> Phosphates and oxalate of calcium. Add excess of ammonia, and then excess of acetic acid; filter.	
	<i>Insoluble.</i> Oxalate of calcium.	<i>Soluble.</i> Phosphates. They may be re-pptd. by ammonia.

Varieties of calculi.—Calculi composed entirely of *uric acid* are common; a minute portion heated on platinum foil chars, burns, and leaves scarcely a trace of ash. The phosphates frequently occur together, forming what is known as the *fusible calculus*, from the readiness with which a fragment aggregates, and even fuses to a bead, when heated on a loop of platinum wire in the blowpipe-flame. The phosphates may, if necessary, be further examined by the method described in connection with urinary deposits. Oxalate of calcium often occurs alone, forming a dark-coloured calculus having a very rough surface, hence termed the *mulberry calculus*. Smaller calculi of the same substance are called, from their appearance, *hempseed calculi*. Calculi of *cystin* are rarely met with. *Xanthin* (from *ξανθός*, *xanthos*, yellow, in allusion to the colour it yields with nitric acid) still less often occurs as a calculus. The earthy concretions or *chalk-stones* which frequently form in the joints of gouty persons are composed chiefly of urates, the sodium salt being that most commonly met with. *Gall-stones* or *biliary calculi* occasionally form in the gall-bladder; they contain *cholesterin* (from *χολή*, *chole*, bile, and *στερεός*, *stereos*, solid), a fatty substance of alcoholoid constitution, soluble in rectified spirit or ether, and crystallizing from such solutions in well-defined, square, scaly

crystals. Phosphatic and other calculi of many pounds weight are occasionally found in the stomach and larger intestines of animals.

QUESTIONS AND EXERCISES.

891. In breathing, how much carbon (in the form of carbonic acid gas) is exhaled from the lungs every 24 hours?

892. How may the presence of carbonic acid gas in expired air be demonstrated?

893. Mention an experiment showing the escape of moisture from the lungs during breathing.

894. State the method of testing for albumen in urine.

895. Give the tests for sugar in urine.

896. What is the average composition of healthy urine?

897. Give the tests for urea.

898. Write the rational formulæ of some compound ureas in which methyl or ethyl displace hydrogen.

899. Describe an artificial process for the production of urea, giving equations.

900. Sketch out a plan for the chemical examination of urinary sediments.

901. A deposit is insoluble in the supernatant urine or in acetic acid; of what substances may it consist?

902. Which compounds are indicated when a deposit redissolves on warming it with the supernatant urine?

903. Name the salts insoluble in warmed urine but dissolved on the addition of acetic acid.

904. Mention the chemical characters of cystin. At what stage of analysis would it be recognised?

905. Describe the microscopical appearance of the following urinary deposits:—

Uric Acid.

Cystin.

Triple phosphate.

Earthy phosphates.

Urates.

Oxalate of Calcium.

Carbonate of Calcium.

Hippuric Acid.

Tube-casts.

Epithelial débris.

Blood.

Pus.

Mucus.

Fat.

Spermatozoa.

Sarcina.

Extraneous Bodies.

906. How are Day's tests for blood, pus, and saliva applied?

907. What is the general, physical, and chemical nature of urinary calculi?

908. How are urinary calculi prepared for chemical examination?

909. Draw out a chart for the chemical examination of urinary calculi?

910. Why is the "fusible calculus" so called? and what is its composition?

911. State the characters of "mulberry" and "hempseed" calculi.

912. What are the "chalk-stones" of gout and "gall-stones" or "biliary calculi"?

THE GALENICAL PREPARATIONS OF THE BRITISH PHARMACOPŒIA.

The preparation of Confections, Decoctions, Enemas, Extracts, Glycerines, Infusions, Inhalations, Juices, Liniments, Lozenges, Mixtures, Ointments, Pills, Plasters, Poultices, Powders, Spirits, Suppositories, Syrups, Tinctures, and Wines, includes a number of mechanical rather than chemical operations, and belongs to the domain of pure Pharmacy. The medical or pharmaceutical pupil will have had ample opportunity of practically studying these compounds before working at experimental chemistry, and will probably have prepared many of them according to the directions of the Pharmacopœia; if not, he is referred to the pages of the last edition of that work for details.

Among the extracts, however, there are five (namely, those of Aconite, Belladonna, Hemlock, Henbane, and Lettuce) which are not simply evaporated infusions, decoctions, or tinctures, like most others, but are evaporated juices from which vegetable albumen, the supposed source of fermentation and decay, has been removed, and chlorophyl (the green colouring-matter of plant-juice) retained practically unim-

paired in tint. For educational practice either of the above-named five raw materials may be employed; but in order that attention may be concentrated on the process by which the extracts are prepared, rather than on any one of the extracts themselves, it suffices to make an extract of some ordinary green vegetable, such as cabbage or turnip-tops. Bruise the green leaves of a good-sized cabbage in a mortar, and press out the juice; heat it gradually to 130° F., and remove the green flocks of chlorophyl which separate, by filtration through calico. When the liquor has all passed through the filter, set the chlorophyl aside for a time, heat the strained liquor to 200° F. to coagulate albumen; remove the latter by filtration and throw it away; evaporate the filtrate by a water-bath to the consistence of thin syrup; then add to it the chlorophyl, and, stirring the whole together assiduously, continue the evaporation at a temperature not exceeding 140° F., until the extract is of a suitable consistence for forming pills. A higher temperature than that indicated would cause the alteration of the chlorophyl to a dark brown substance, any such extract used in pharmacy no longer having the green tint which custom and the British Pharmacopœia demand.

QUESTIONS AND EXERCISES.

913. Enumerate the different classes into which official galenical preparations may be divided.

914. Describe the general process for the preparation of green extracts:—

Aconite.

Hemlock.

Belladonna.

Henbane.

Lettuce.

915. Why is vegetable albumen excluded in the preparation of green extracts?

916. How may chlorophyl be removed from vegetable juices, and again be introduced into their evaporated residues, without destroying its colour?

917. For what reason is exposure of chlorophyl to a boiling temperature avoided in the manufacture of green extracts?

THE CHEMICAL PREPARATIONS OF THE BRITISH PHARMACOPŒIA.

The process by which every official chemical substance is prepared has already been described, and the strict chemical character of the processes illustrated by experiments and explained by aid of equations. Should the reader, in addition, desire an intimate acquaintance with those details of manipulation on which the successful and economic manufacture of chemical substances depends, he is advised to prepare, if he has not done so already, a few ounces of each of the salts mentioned in the British Pharmacopœia or commonly used in Pharmacy. An additional guide in these operations will be the Pharmacopœia itself.

The production of many chemical and galenical substances on a commercial scale can only be successfully carried on in manufacturing laboratories, and with some knowledge of the circumstances of supply and demand, value of raw material, and of by-products, etc.; for the technical preparation of such substances requires much knowledge beyond even a thorough acquaintance with chemistry. Still, in the present day, commercial Chemistry and Pharmacy can best hope for success when founded on the working out of abstract scientific principles. The problem of manufacturing success is now only solved with certainty by sound and wisely-applied science.

Memorandum.—The next subjects of experimental study will be determined by the nature of the student's future pursuits. In most cases the operations of quantitative analysis will engage attention. These should be of a volumetric and gravimetric character; for details concerning them see the following pages.

QUANTITATIVE ANALYSIS.

INTRODUCTORY REMARKS.

General principles.—The proportions in which chemical substances unite with each other in forming compounds are definite and invariable (p. 44). Quantitative analysis is based on this law. When, for example, aqueous solutions of a salt of silver and a chloride are mixed, a white curdy precipitate is produced containing chlorine and silver in atomic proportions—that is, 35·5 parts of chlorine to 108 of silver. No matter what the chloride or what the salt of silver, the resulting chloride of silver is invariable in composition. The formula AgCl is a convenient picture of this compound in these proportions. The weight of a definite compound being given, therefore, the proportional amounts of its constituents can be ascertained by simple calculation. Suppose, for instance, 8·53 parts of chloride of silver have been obtained in some analytical operation: this amount will contain 2·11 parts of chlorine and 6·42 of silver. For if 143·5 (the molecular weight) of chloride of silver contain 35·5 (the atomic weight) of chlorine, 8·53 of chloride of silver will be found to contain 2·11 of chlorine:—

$$\begin{array}{rcl}
 143\cdot5 & : & 35\cdot5 \\
 & & 8\cdot53 \\
 \hline
 & & 1\cdot065 \\
 & & 17\cdot75 \\
 & & 284\cdot0 \\
 \hline
 143\cdot5) & 2815(2\cdot11 & \\
 & 287\cdot0 & \\
 \hline
 & 15\cdot81 & \\
 & 14\cdot35 & \\
 \hline
 & 1\cdot465 & \\
 & 1\cdot435 & \\
 \hline
 & & x = 2\cdot11
 \end{array}$$

And if 143·5 of chloride of silver contain 108 of silver, 8·53 of chloride of silver will contain 6·42 of silver. To ascertain, for

example, the amount of silver in a substance, containing, say, nitrate of silver, all that is necessary is to take a weighed quantity of the substance, dissolve it, precipitate the whole of the silver by adding hydrochloric acid or other chloride till no more chloride of silver falls, collect the precipitate on a filter, wash, dry, and weigh. The amount of silver in the dried chloride, ascertained by calculation, is the amount of silver in the quantity of substance on which the operation was conducted; a rule-of-three sum gives the quantity per cent.—the form in which the results of quantitative analysis are usually stated. Occasionally a constituent of a substance admits of being isolated and weighed in the uncombined state. Thus the amount of mercury in a substance may be determined by separating and weighing the mercury in the metallic condition; if occurring as calomel (HgCl) or corrosive sublimate (HgCl_2), the proportion of chlorine may then be ascertained by calculation ($\text{Hg}=200$; $\text{Cl}=35.5$).

Nature of Gravimetric Quantitative Analysis.—As above stated, a body may be isolated and weighed and its quantity thus ascertained; or it may be separated and weighed in combination with another body whose combining proportion is well known; this is quantitative analysis by the *gravimetric* method.

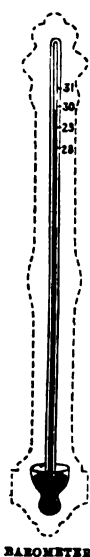
Nature of Volumetric Quantitative Analysis.—Volumetric operations depend for success on some accurate initial gravimetric operation. A weighed amount of a pure salt is dissolved in a given volume of water or other fluid and thus forms a *standard solution*. Accurately measured quantities of such a solution will obviously contain just as definite amounts of the dissolved salt as if those amounts were actually weighed in a balance, and as measuring occupies less time than weighing, the volumetric operations can be conducted with great economy of time as compared with the corresponding gravimetric operations. Quantitative analysis by the *volumetric* method consists in noting the volume of the standard liquid required to be added to the substance under examination before a given effect is produced. Thus, for instance, a solution of nitrate of silver of known strength may be used in experimentally ascertaining an unknown amount of a chloride in any substance. The silver solution is added to a solution of a definite quantity of the substance until flocks of chloride of silver cease to be precipitated: every 108 parts of silver added (or 170 of nitrate of silver: $\text{Ag}=108$, $\text{N}=14$, $\text{O}_3=48$; total 170) indicates the

presence of 35.5 of chlorine, or an equivalent quantity of any chloride. The preparation of standard solutions, such as that of the nitrate of silver, to which allusion is here made, requires considerable care; but when made, certain analyses can, as already indicated, be executed with far more rapidity and ease than by gravimetric processes.

Quantitative determination of (a) atmospheric pressure, (b) temperature, and (c) weight.—The quantitative analysis of solids and liquids often involves quantitative determinations of atmospheric pressure, temperature, and weight. These processes will now be explained, after which an outline of volumetric and gravimetric quantitative analysis will be given. The scope of this work precludes any attempt to describe all the little mechanical details observed by quantitative analysts; essential operations, however, are so fully treated that expert manipulators will meet with little difficulty.

QUANTITATIVE DETERMINATION OF ATMOSPHERIC PRESSURE.

Fig. 62.

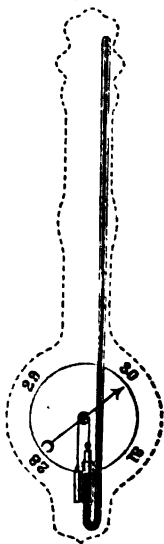


BAROMETER.

The Barometer.—The analysis of gases and vapours, especially involves determinations of the varying pressure of the atmosphere as indicated by the barometer (from *βάρος*, *baros*, weight, and *μέτρον*, *metron*, measure).

The ordinary mercurial barometer is a glass tube 33 or 34 inches long, closed at one end, filled with mercury, and inverted in a small cistern or cup of mercury (fig. 62). The mercury remains in the tube owing to the weight or pressure of the atmosphere on the exposed surface of the liquid, the average height of the column being nearly 30 inches. In the popular form of the instrument, the wheel-barometer, the cistern is formed by a recurvature of the tube (fig. 63); on the exposed surface of the mercury a float is placed, from which a

Fig. 63.



BAROMETER.

thread passes over a pulley and moves an index whenever the column of mercury rises or falls. As supplied to the public, these barometers are usually enclosed in ornamental frames with thermometers attached. In the wheel-barometer the glass tube and contained column of mercury are altogether enclosed, the index alone being visible. In the other variety the upper end of the glass tube and mercurial column are exposed, and the height of the mercury is ascertained by direct observation.

The aneroid barometer (from *ἀ, a*, without, and *νηρός, nēros*, fluid) consists of a small shallow vacuous metal drum, the sides of which approach each other when an increase of atmospheric pressure occurs, their elasticity enabling them to recede towards their former position on a decrease of pressure. This motion is so multiplied and altered in direction by levers, etc., as to act on a hand traversing a plate on which are marked numbers corresponding with those showing the height of the mercurial column of the ordinary barometer by which the aneroid was adjusted. *The Bourdon barometer* (from the name of the inventor) is a modified aneroid, containing in the place of the round metal box a flattened vacuous tube of metal bent nearly to a circle. These barometers are also useful for measuring the pressure in steam-boilers, etc. Under the name of *pressure-gauges* they are sold to indicate pressures of 500 pounds per square inch and upwards. From their portability (they can be made of 1 to 2 inches in diameter and 1 inch thick) they are excellent companions for travellers wishing to know the height of hills, mountains, and other elevations.

For further information concerning the influence of pressure on the volume of a gas or vapour *see* p. 635; and for descriptions of the methods of analysing gases, refer to Ganot's "Physics" (translated by Atkinson), Miller's "Chemical Physics," and "Analysis of Gases" in Watts's "Dictionary of Chemistry."

QUANTITATIVE DETERMINATION OF TEMPERATURE.

General Principles.—As a rule, all bodies expand on the addition, and contract on the abstraction of heat, the alteration in volume being constant and regular for equal increments or decrements of temperature. The extent of this alteration in a given substance, expressed in parts or degrees, constitutes the usual method of intelligibly stating with accuracy, precision,

and minuteness, a particular condition of warmth or temperature—that is, of sensible heat. The substance commonly employed for this purpose is mercury, the chief advantages of which are that it will bear a high temperature without boiling, a low temperature without freezing, does not adhere to glass to a sufficient extent to “wet” the sides of any tube in which it may be enclosed, and, from its good conducting-power for heat, responds rapidly to changes of temperature. Platinum, earthenware, alcohol, and air are also occasionally used for thermometric purposes.

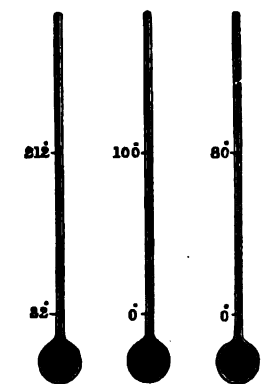
The Thermometer.—The construction of an accurate thermometer is a matter of great difficulty; but the following are the leading steps in the operation. Select a piece of glass tubing having a fine capillary (*capillus*, a hair) bore, and about a foot long; heat one extremity in the blowpipe-flame until the orifice closes, and the glass is sufficiently soft to admit of a bulb being blown; heat the bulb to expel air, immediately plunging the open extremity of the tube into mercury; the bulb having cooled, and some mercury having entered and taken the place of expelled air, again heat the bulb and tube until the mercury boils and its vapour escapes through the bore of the tube; again plunge the extremity under mercury, which will probably now completely fill the bulb and tube. When cold the bulb is placed in melting ice. The top of the column of mercury in the capillary tube should then be within an inch or two of the bulb; if higher, some of the mercury must be expelled by heat; if lower, more metal must be introduced as before. The tube is now heated near the open end and a portion drawn out, until the diameter is reduced to about one tenth. The bulb is next warmed until the mercurial column rises above the constricted part of the tube, which is then rapidly fused in the blowpipe-flame, and the extremity of the tube removed.

The instrument is now ready for *graduation*. The bulb is placed in boiling water (a medium having, *cæteris paribus*, an invariable temperature), and, when the position of the top of the mercurial column is constant, a mark is made on the tube by a scratching diamond or a file. This operation is repeated

with melting ice (also a medium having an invariable temperature). The space between these two marks is divided into a certain number of intervals termed *degrees*. Unfortunately this number is not uniform in all countries: in England it is 180, as proposed by Fahrenheit; in France 100, as proposed by Celsius (the Centigrade scale), a number generally adopted by scientific men: in some parts of the Continent the divisions are 80 for the same interval, as suggested by Réaumur. Whichever be the number selected, similar markings should be continued beyond the boiling- and freezing-points as far as the length of the stem admits. They may be made on the stem itself, or on any wood, metal, or earthenware frame on which the stem is mounted.

Fig. 64.

THERMOMETRIC SCALES.



Fahrenheit. Centigrade. Réaumur.

Thermometric Scales (fig. 64).—On the Centigrade (C.) and Réaumur (R.) scales the freezing point of water is made zero, and the boiling-point 100 and 80 respectively; on the Fahrenheit (F.) scale the zero is placed 32 degrees below the congealing-point of water, the boiling-point of which becomes, consequently, 212. Even on the Fahrenheit system temperatures below the freezing-point of water are often spoken of as “degrees of frost;” thus 19° as marked on the thermometer

would be regarded as "13 degrees of frost." It is to be regretted that the freezing-point of water is not universally regarded as the zero-point, and the number of intervals between that and the boiling-point everywhere the same.

The degrees of one scale are easily converted into those of another if their relations be remembered, namely:—180 (F.), 100 (C.), 80 (R.); or 18, 10, and 8; or best, 9, 5, and 4.

Formulae for the conversion of degrees of one thermometric scale into those of another.

F=Fahrenheit.
R=Réaumur.

C=Centigrade.
D=The observed degree.

If above the freezing-point of water (32° F; 0° C; 0° R),

F into C		$(D - 32) \div 9 \times 5.$
F " R		$(D - 32) \div 9 \times 4.$
C " F		$D \div 5 \times 9 + 32.$
R " F		$D \div 4 \times 9 + 32.$

If below freezing, but above 0° F (−17·77° C; −14·22° R),

F into C		$-(32 - D) \div 9 \times 5.$
F " R		$-(32 - D) \div 9 \times 4.$
C " F		$32 - (D \div 5 \times 9).$
R " F		$32 - (D \div 4 \times 9).$

If below 0° F (−17·77° C; −14·22° R),

F into C		$-(D + 32) \div 9 \times 5.$
F " R		$-(D + 32) \div 9 \times 4.$
C " F		$-(D \div 5 \times 9) - 32.$
R " F		$-(D \div 4 \times 9) - 32.$

For all degrees:—

C into R		$D \div 5 \times 4.$
R " C		$D \div 4 \times 5.$

In ascertaining the temperature of a liquid, the bulb of a thermometer is simply inserted and the degree noted. In determining the boiling-point, also, the bulb is inserted in the liquid, if a pure substance. In taking the boiling-point of a liquid which is being distilled from a mixture, the bulb of the thermometer should be near to but not beneath the surface.

The "boiling-point" of a liquid is the temperature at which the elasticity of the vapour of the substance overcomes the atmospheric or other pressure to which the liquid is exposed.

If the pressure is equal to 760 mm. (29·92 inches) of mercury, water will boil at 100° C. (212° F.). The boiling-point of a drop of a fluid is taken by introducing it into the closed extremity of a small U tube, the remaining portion of the closed limb being filled with mercury. The tube is lowered into a bath, the open limb being above the surface of the fluid of the bath. The bath is slowly and equally heated, and the boiling-point of the liquid, indicated by the mercury falling until it is level in the two limbs, taken by a thermometer whose bulb is close to the U tube.

The following are the boiling-points of a few substances met with in pharmacy:—

	Centigrade.	Fahrenheit.
Alcohol, absolute	78·3	173
„ 84 per cent.	79·5	175
„ 49 per cent. (proof spirit)	81·4	178·5
„ amylie	132·2	270
Benzol	80·6	177
Bromine	63	145·4
Benzoic acid	239·0	462
Carbolic acid	187·8	370
Chloroform	61	142
Ether (B. P.) (below)	40·5	105
„ pure	35	95
Mercury <i>in vacuo</i> (as in a thermometer) .	304	580
„ in air (barom. at 30 inches) . .	350	662
Water (barom. at 29·92 inches)	100	212
„ („ 29·33 „)	99·5	211
„ („ 28·74 „)	99	210
Saturated solutions of:—		
Cream of tartar	101	214
Common salt	106·6	224
Sal-ammoniac	113·3	236
Nitrate of sodium	119	246
Acetate of sodium	124·4	256
Chloride of calcium	179·4	355

To determine melting-points of fats.—Heat a fragment of the substance (spermaceti or wax for example) till it liquefies, and then draw up a small portion into a thin glass tube, about

the size of a knitting-needle. Immerse the tube in cold water contained in a beaker, and slowly heat the vessel till the thin opaque cylinder of solid fat melts and becomes transparent; a delicate thermometer placed in the water indicates the point of change to the fifth of a degree. Remove the source of heat, and note the congealing-point of the substance; it will, in nearly all cases, be identical with or close to the melting-point.

Pyrometers.—Temperatures above the boiling-point of mercury are determined by ascertaining to what extent a bar of platinum or porcelain has elongated. The bar is enclosed in a cavity of a suitable case, a plug of platinum or porcelain placed at one end of the bar, and the whole exposed in the region the temperature of which is to be found. After cooling, the distance to which the bar has forced the plug along the cavity is accurately measured and the corresponding degree of temperature noted. The value of the distance is fixed for low temperatures by comparison with a mercurial thermometer, and the scale carried upwards through intervals of equivalent length. Such thermometers are conventionally distinguished from ordinary instruments by the name *pyrometer* (from $\pi\upsilon\rho$, *pur*, fire, and $\mu\epsilon\tau\rho\omicron\nu$, *metron*, measure).

The following are melting-points of substances official in the British Pharmacopœia :—

	In degrees Centigrade.	In degrees Fahrenheit.
Acetic acid, glacial	8·9	48
" " congeals at	1·1	34
Benzoic acid	120	248
Carbolic acid	35	95
Oil of theobroma (about)	32	90
Phosphorus	43·3	110
Prepared lard (about)	38	100
" suet	39·5	103
Spermaceti (not under)	38	100
White wax "	65·5	150
Yellow wax "	60	140

The order of fusibility of a few of the metals is as follows:—

	In degrees Centigrade.	In degrees Fahrenheit.
Mercury	— 39·4	— 39
Potassium	+ 62·5	+144·5
Sodium	97·6	207·7
Tin	227·8	442
Bismuth	264	507
Lead	325	617
Zinc	411·6	773
Antimony	621	1150
Silver	1023	1873
Copper	1091	1996
Gold	1102	2016
Cast iron	1530	2786

QUESTIONS AND EXERCISES.

918. On what fundamental laws are the operations of quantitative analysis based?

919. What is the general nature of *gravimetric* quantitative analysis?

920. Describe the general principle of *volumetric* quantitative analysis.

921. How are variations in atmospheric pressure quantitatively determined?

922. Explain the construction and mode of action of a mercurial barometer.

923. In what respect does a wheel-barometer differ from an instrument in which the readings are taken from the top of the column of mercury?

924. Describe the principle of action of an aneroid barometer.

925. On what general principles are thermometers constructed?

926. What material is employed in making thermometers?

927. Why is mercury selected as a thermometric indicator?

928. Describe the manufacture of a mercurial thermometer.

929. How are thermometers graduated?

930. Give formulæ for the conversion of the degrees of one thermometric scale into those of another, (a) when the temperature is above the freezing-point of water, (b) below 32° F. but above 0° F., and (c) below 0° F.

931. Name the degree C. equivalent to 60° F.

932. What degree C. is represented by -4° F.?

933. Mention the degree F. indicated by 20° C.

934. Convert 100° R. into degrees C. and F.

935. State the boiling-points of alcohol, chloroform, ether, mercury, and water on either thermometric scale.

936. Describe the details of manipulation in estimating the melting-points of fats.

937. In what respect do pyrometers differ from thermometers?

938. Mention the melting-points of glacial acetic acid, oil of theobroma, lard, suet, and wax.

939. Give the fusing-points of tin, lead, zinc, copper, and cast iron.

QUANTITATIVE DETERMINATION OF WEIGHT.

DEFINITIONS.

All bodies, celestial and terrestrial, attract each other, the amount of attraction being in direct proportion to the quantity of matter of which they consist, and in inverse proportion to the squares of their distances. This is *gravitation*. When gravitation in certain directions is exactly counterbalanced by gravitation in opposite directions, a body (e.g. the earth) remains suspended in space. Such a body, in relation to other bodies, has gravity but not weight. *Weight* is the effect of gravity, being the excess of gravitation in one direction over and above that exerted in the opposite direction. Weight, truly, in any terrestrial substance, is the excess of attraction which it and the earth have for each other over and above the attraction of each in opposite directions by the various heavenly bodies. But, practically, the weight of any terrestrial substance is the effect of the attraction of the earth only. *Specific weight* is the definite or precise weight of a body in relation to its bulk; it is more usually but not quite correctly termed *specific gravity*—gravity belonging to the earth, and not, in any sensible degree, to the substance.

QUESTIONS AND EXERCISES.

940. What is understood by gravitation ?
941. State the difference between weight and gravity.
942. Mention a case in which a body has gravity but no apparent weight.
943. Practically, what causes the weight of terrestrial substances ?

Weights and Measures.

The Balance.—The balance used in the quantitative operations of analytical chemistry must be accurate and sensitive. The points of suspension of the beam and pan should be polished steel or agate knife-edges, working on agate planes. It should turn easily and quickly, without too much oscillation, to $\frac{1}{800}$ or $\frac{1}{600}$ of a grain, or $\frac{1}{10}$ of a milligramme, when 1000 grains, or 50 or 60 grammes, are placed in each pan. (Grammes are weights of the metric system, a description of which is given on the next two or three pages.) The beam should be light but strong, capable of supporting a load of 1500 grains or 100 grammes; its oscillations are observed by help of a long index attached to its centre, and continued downwards for some distance in front of the supporting pillar of the balance. The instrument should be provided with screws for purposes of adjustment, a mechanical contrivance for supporting the beam above its bearings when not in use or during the removal or addition of weights, spirit-levels to enable the operator to give it a horizontal position, and be enclosed in a glass case to protect it from dust. It should be placed in a room the atmosphere of which is not liable to be contaminated by acid fumes, in a situation free from vibration; and a vessel containing lumps of quicklime should be placed in the case to keep the enclosed air dry and prevent the formation of rust on any steel knife-edges or other parts. During weighing, the doors of the balance should be shut, in order that currents of air may not unequally influence the pans.

The Weights.—These should be preserved in a box having a separate compartment for each. They must not be lifted directly with the fingers, but by a small pair of forceps. If

grain-weights, they should range from 1000 gr. to $\frac{1}{10}$ gr., a $\frac{1}{10}$ weight being fashioned of gold wire to act as a "rider" on the divided beam, and thus indicate by its position 100ths and 1000ths of a grain. From $\frac{1}{10}$ to 10 gra. the weights may be of platinum; thence upwards to 1000 gra. of brass. The relation of the weights to each other should be decimal. Metric decimal weights may range from 100 grammes to 1 gramme, of brass, and thence downwards to 1 centigramme, of platinum, a gold centigramme rider being employed to indicate milligrammes and tenths of a milligramme.

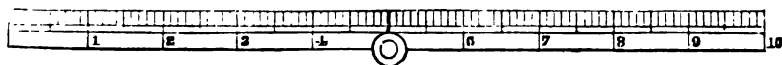
The Metric System of weights (the word *metric* is from the Greek μέτρον, *metron*, measure) is greatly to be preferred to the British, the relation of the metric weights of all denominations to measures of length, capacity, and surface being so simple as to be within the perfect comprehension of a child; while under the British plan, the weights have no such relation, either with each other or with the various measures. Moreover, the metric system is in perfect harmony with the universal method of counting; it is a decimal system.

[It is perhaps impossible to realize, much more express, the advantages we enjoy from the fact that in every country of the world the system of numeration is identical. That system is the decimal. Whatever language a man speaks, his method of numbering is decimal; his talk concerning numbers is decimal; his written or printed signs signifying number are decimal. With the figures 1, 2, 3, 4, 5, 6, 7, 8, 9, 0 he represents all possible variation in number, the position of a figure in reference to its companions alone determining its value, a figure on the left hand of any other figure in an allocation of numeral symbols (for example, 1869) having ten times the value of that figure, while the figure on the right hand of any other has a tenth of the value of that other. When the youngest pupil is asked how many units there are in 1869, he smiles at the simplicity of the question, and says 1869. How many tens? 186, and 9 over. How many hundreds? 18, and 69 over. How many thousands? 1, and 869 over. But if he is asked how many scruples there are in 1869 grains, how many drachms, how many ounces—he brings out his slate and pencil. And so with the pints or gallons in 1869 fluid ounces, or the feet and yards in 1869 inches, or the pence, shillings, and pounds in 1869 farthings; to say nothing of cross questions, such as the value of 1869 articles at 9s. 6d. per dozen,

or of the perplexity caused by the varying values of several individual weights or of measures of length, capacity, and surface in different parts of the country. What is desired is, that there should be an equally simple decimal relation among weights and measures and coins as already universally exists among numbers. This condition of things having already been introduced into most other countries, there is no good reason why it should not be accomplished in our own.]

The system of weights and measures legalized by "the Metric Weights and Measures Act, 1864," is founded on the *mètre*. Fig. 65 represents a pocket folding measure the tenth part of a *mètre* in length, divided into 10 centimètres, and each centimètre into 10 millimètres.

Fig. 65.



THE DÉCIMÈTRE.

The units of the system with their multiples and submultiples are as follows:—

UNITS.

Length.—The *Unit of Length* is the *MÈTRE*, derived from the measurement of the Quadrant of a Meridian of the Earth. (Practically, it is the length of certain carefully preserved bars of metal, from which copies have been taken.)

Surface.—The *Unit of Surface* is the *ARE*, which is the Square of Ten *Mètres*.

Capacity.—The *Unit of Capacity* is the *LITRE*, which is the Cube of a Tenth part of a *Mètre*.

Weight.—The *Unit of Weight* is the *GRAMME*, which is the Weight of that quantity of distilled water, at its maximum density, which fills a Cube of the One hundredth part of the *Mètre*.

TABLE.

Note.—Multiples are denoted by the Greek words "Deka," Ten, "Hecto," Hundred, "Kilo," Thousand.

Subdivisions by the Latin words, "Deci," One tenth, "Centi," One hundredth, "Milli," One thousandth.

Quantities.	Length.	Surface.	Capacity.	Weight.
1000	Kilo-mètre	...	Kilo-litre	Kilo-gramme
100	Hecto-mètre	Hectare	Hecto-litre	Hecto-gramme
10	Déca-mètre	...	Déca-litre	Déca-gramme
1 (Units)	MÈTRE	ARE	LITRE	GRAMME
·1	Déci-mètre	...	Déci-litre	Déci-gramme
·01	Centi-mètre	Centiare	Centi-litre	Centi-gramme
·001	Milli-mètre	...	Milli-litre	Milli-gramme

When the metric method is exclusively adopted in Great Britain, these Units and Table, comprising the entire System of Weights and Measures, represent all that will be essential to be learnt in lieu of the numerous and complicated Tables hitherto in use. Adopting the style of elementary books on arithmetic, the Table may be expanded in the following manner :—

10 Milligrammes	make 1 centigramme.
10 Centigrammes	„ 1 décigramme.
10 Décigrammes	„ 1 gramme.
10 Grammes	„ 1 décagramme.
10 Décagrammes	„ 1 hectogramme.
10 Hectogrammes	„ 1 kilogramme.

10 Millilitres make 1 centilitre,
etc.

10 Millimètres make 1 centimètre,
etc.

The following approximate British equivalents of metrical units should be committed to memory :—

1 Mètre	= 3 feet 3 inches and 3 eighths.
1 Are	= a square whose side is 11 yards.
1 Litre	= 1½ pint.
1 Gramme	= 15½ grains.

The Kilomètre is equal to 1100 yards.

The Hectare = 2½ acres nearly.

The Metric Ton of 1000 Kilogrammes = 19 cwt. 2 qrs. 20 lbs. 10 oz.

The Kilogramme = 2 lbs. 3½ oz. nearly.

For *exact* equivalents, in many forms, *see* pp. 588 and 589. (The word *gramme* is sometimes, unfortunately, written *gram*.)

Decimal Coinage.—In most countries where the metric system of weights and measures is employed, a decimal division of coins is also adopted. This course, conjoined with the ordinary decimal method of enumerating, which fortun-

ately is in universal use, renders calculations of all kinds most simple,—easy to an extent which cannot be conceived in countries like England, where the operations of weighing, measuring, paying, and counting have only the most absurdly intricate relations to each other.

The General Council under whose authority the British Pharmacopœia is issued encourages medical practitioners and pharmacists in the adoption of the metric system, and gives the annexed statement of metric weights and measures.

WEIGHTS AND MEASURES OF THE METRICAL SYSTEM.

(From the British Pharmacopœia of 1867.)

WEIGHTS.

1 Milligramme	= the thousandth part of one grm. or 0·001 grm.	
1 Centigramme	= the hundredth	0·01 "
1 Décigramme	= the tenth	0·1 "
1 Gramme	= weight of a cubic centimètre of water at 4° C.	1·0 "
1 Décagramme	= ten grammes	10·0 "
1 Hectogramme	= one hundred grammes	100·0 "
1 Kilogramme	= one thousand grammes	1000·0 (1 kilo.).

MEASURES OF CAPACITY.

1 Millilitre	=	1 cub. centim. or the mea. of 1 gram. of water.	
1 Centilitre	=	10	10 " "
1 Décilitre	=	100	100 " "
1 Litre	=	1000	1000 " (1 kilo.).

MEASURES OF LENGTH.

1 Millimètre	= the thousandth part of one mètre or 0·001 mètre.	
1 Centimètre	= the hundredth	0·01 "
1 Décimètre	= the tenth	0·1 "
1 Mètre	= the ten-millionth part of a quarter of the meridian of the earth.	

METRICAL MEASURES OF LENGTH.

	In English inches.	In English feet =12 inches.	In English yards =3 feet.	In English fathoms =6 feet.	In English miles =1760 yards.
Millimètre	0.03937	0.003281	0.0010936	0.0005468	0.0000006
Centimètre	0.39371	0.032809	0.0109363	0.0054682	0.0000062
Décimètre	3.93708	0.328030	0.1093633	0.0546816	0.0000621
Mètre	39.37079	3.280899	1.0936331	0.5468165	0.0006214
Décamètre	393.70790	32.808992	10.9363306	5.4681653	0.0062138
Hectomètre	3937.07900	328.089917	109.3633056	54.6816528	0.0621382
Kilomètre	39370.79000	3280.899167	1093.6330556	546.8165278	0.6213824
Myriomètre	393707.90000	32808.991667	10936.3305556	5468.1652778	6.2138242

1 inch = 2.539954 centimètres.

1 foot = 3.0479449 décimètres.

1 yard = 0.9143835 mètre.

1 mile = 1.6093149 kilomètre.

METRICAL MEASURES OF SURFACE.

	In English square feet.	In English sq. yards =9 square feet.	In English poles= 27.25 square feet.	In English rods= 10890 square feet.	In English acres =43560 sq. feet.
Centiare or square mètre .	10.764399	1.196033	0.0395383	0.0009885	0.0002471
Are or 100 square mètres .	1076.429934	119.603326	3.9538290	0.0988457	0.0247114
Hectare or 10,000 sq. mètres	107642.993418	11960.332602	395.3828959	9.8845724	2.4711431

1 square inch = 6.4513669 square centimètres.

1 square foot = 9.2903384 square décimètres.

1 square yard = 0.83604715 square mètre or centiare.

1 acre = 0.40467102 hectare.

1 square mile = 2.58998451 square kilomètre.

METRICAL MEASURES OF CAPACITY.

	In cubic inches.	In cubic feet= 1728 cubic inches.	In pints= 34.6633 cubic inches.	In gallons= 8 pints=277.27364 cubic inches.	In bushels= 8 gal- lons=231.9072 cubic inches.
Millilitre, or cubic centimètre	0.06103	0.000035	0.00176	0.0002201	0.0000275
Centilitre, or 10 cu. centimètres	0.61027	0.000353	0.01761	0.0022010	0.0002751
Déclilitre, or 100 cu. centimètres	6.10271	0.003532	0.17608	0.0220097	0.0027512
Litre, or cubic décimètre . .	61.02705	0.035317	1.76077	0.2200967	0.0275121
Décilitre, or centistère . .	610.27052	0.353166	17.60773	2.2009668	0.2751208
Hectolitre, or décastère . .	6102.70515	3.531658	176.07734	22.0096677	2.7512085
Kilolitre, or stère, or cu. mètre	61027.05152	35.316581	1760.77341	220.0966767	27.5120846
Myriolitre, or décastère . .	610270.51519	353.165807	17607.73414	2200.9667675	275.1208459
1 cubic inch = 16.896176 cubic centimètres. 1 cubic foot = 28.315812 cubic décimètres. 1 gallon = 4.535986 litres.					

METRICAL MEASURES OF WEIGHT.

	In English grains.	In troy ounces =480 grains.	In avoirdupois lbs. =7000 grains.	In cwt.=112 lbs. =784000 grains.	Tons=20 cwt.= 15680000 grains.
Milligramme	0.01543	0.000032	0.0000022	0.0000000	0.0000000
Centigramme	0.15432	0.000022	0.0000220	0.0000002	0.0000000
Déigramme	1.54323	0.003215	0.0002205	0.0000020	0.0000001
Gramme	15.43235	0.032151	0.0022046	0.0000197	0.0000010
Déagramme	154.32349	0.321507	0.0220462	0.0001968	0.0000098
Hectogramme	1543.23488	3.215073	0.2204621	0.0019684	0.0000984
Kilogramme	15432.34880	32.150727	2.2046213	0.0196841	0.0009842
Myriogramme	154323.48800	321.507267	22.0462126	0.1968412	0.0098421
1 grain = 0.064799 gramme. 1 troy oz. = 31.103496 grams. 1 lb. avd. = 0.453598 kilogram. 1 cwt. = 50.802377 kilograms.					

The foregoing Tables, from the Diary of Messrs. De La Rue, and the following from the British Pharmacopœia, will be found useful for reference :—

WEIGHTS AND MEASURES OF THE BRITISH PHARMACOPŒIA OF 1867.

WEIGHTS.

1 Grain	gr.		
1 Ounce	oz.	=	437·5 grains.
1 Pound	lb.	= 16 ounces	= 7000 „

MEASURES OF CAPACITY.

1 Minim	min.		
1 Fluid Drachm	fl. drm.	=	60 minims.
1 Fluid Ounce	fl. oz.	=	8 fluid drachms.
1 Pint	O.	=	20 fluid ounces.
1 Gallon	C.	=	8 pints.

MEASURES OF LENGTH.

1 line = $\frac{1}{12}$ inch.

1 inch = $\frac{1}{39 \cdot 1393}$ seconds-pendulum.

12 „ = 1 foot.

36 „ = 3 feet = 1 yard.

Length of pendulum vibrating seconds of mean }
time in the latitude of London, in a vacuum } 39·1393 inches.
at the level of the sea }

(1 cubic inch of distilled water at 62° F. and 30 inch Barom. = 252·458 grains.)

RELATION OF MEASURES TO WEIGHTS.

1 Minim is the measure of		0·91 grain of water.
1 Fluid Drachm „		54·68 grains of water.
1 Fluid Ounce „	1 ounce or	437·5 „
1 Pint „	1·25 pound or	8750·0 „
1 Gallon „	10 pounds or	70,000·0 „

(Gtt. = *guttae*, *drops*. The term “drop” indicates a quantity which is indefinite, and should only be used when approximateness is alone desired.)

QUESTIONS AND EXERCISES.

944. Mention some advantages of a decimal system of weights and measures.

945. What is the name of the chief unit of the metric decimal system of weights and measures?

946. Mention the names of the metric units of surface, capacity, and weight, and state how they are derived from the unit of length.

947. How are multiples of metric units indicated?

948. State the designations of submultiples of metric units.

949. How many mètres are there in a kilomètre?

950. How many millimètres in a mètre?

951. How many grammes in 5 kilogrammes?

952. How many milligrammes in $13\frac{1}{2}$ grammes?

953. In 1869 centigrammes how many grammes?

954. In a mètre measure 5 centimètres wide and 1 centimètre thick how many cubic centimètres?

955. How many litres are contained in a cubic mètre of any liquid?

956. State the British equivalent of the mètre.

957. How many square yards in an are?

958. How many fluid ounces in a litre?

959. How many ounces in a kilogramme?

960. Give the relation of a metric ton (1000 kilos.) to a British ton.

961. How many grains are there in 1 British ton?

962. How many ounces in 1 ton?

963. How many grains of water in 1 fluid drachm?

964. How many minims in 1 pint?

965. How many grains in 1 pint of water?

966. Whence is the British unit of length derived?

Specific Weight or Specific Gravity.

The *specific* weight of a substance is its weight in comparison with the weights of similar bulks of other substances. This comparative heaviness of *solids and liquids* is conventionally expressed in relation to water: they are considered as being lighter or heavier than water. Thus, water being regarded as unity = 1, the relative weight, or specific weight,

of ether is represented by the figures .720 (it is nearly three-fourths, .750, the weight of water), oil of vitriol by 1.843 (it is nearly twice, 2.000, as heavy as water). The specific weight of substances is, moreover, the weight of similar volumes at *sixty degrees* (60° F.); for the weight of a definite volume of any substance will vary according to temperature, becoming heavier when cooled and lighter when heated, different bodies (gases excepted) differing in their rate of contraction and expansion. While, then, specific weight or, conventionally, *specific gravity* is, truly, the comparative weight of equal bulks, the numbers which, in Great Britain, commonly represent specific gravities are the comparative weights of equal bulks at 60° F., water being taken as unity.* The standard of comparison for gases was formerly air, but is now usually hydrogen.

SPECIFIC GRAVITY OF LIQUIDS.

Procure any small bottle holding from 100 to 1000 grains (fig. 66), and having a narrow neck; counterpoise it in a delicate balance; fill it to about halfway up the neck with pure distilled water having a temperature of 60° F.; ascertain the weight of the water, and, for convenience, add or subtract a drop or two, so that the weight shall be a round number of grains; mark the neck by a diamond or file-point at the part cut by the lower edge of the curved surface of the water. Consecutively fill up the bottle to the neck-mark with several other liquids, cooled or warmed to 60° F., first rinsing out the bottle once or twice with a small quantity of each liquid, and note the weights; the respective figures will represent the relative weights of equal bulks of the liquids. If the capacity of the bottle is 10, 100, or 1000 grains, the resulting weights

* The true weight of a body is its weight in air plus the weight of an equal bulk of air and minus the weight of a bulk of air equal to the bulk of the brass or other weights employed; or, in other words, its weight *in vacuo*, uninfluenced by the buoyancy of the air: but such a correction of the weight of a body is seldom necessary, or, indeed, desirable. *Density* is sometimes improperly regarded as synonymous with *specific gravity*. It is true that the density of a body is in exact proportion to its specific gravity; but the former is more correctly the comparative bulk of equal weights, while specific gravity is the comparative weight of equal bulks.

will, without calculation, show the specific gravities of the liquids; if any other number a rule-of-three sum must be worked out to ascertain the weight of the liquids as compared with 1 (or 1·000) of water. Bottles conveniently adjusted

Fig. 66.

Fig. 67.

Fig. 68.

Fig. 69.



SPECIFIC-GRAVITY BOTTLES.

to contain 250, 500, or 1000 grains, or 100 or 50 grammes of water, when filled to the top of their perforated stopper (fig. 68), and other forms of the instrument (fig. 67 and 69), are sold by all chemical-apparatus makers. Figure 69 is that of a bottle extremely useful in ascertaining the specific gravities of very volatile liquids.

Verify some of the following stated specific gravities of official liquids:—

Acid, acetic	1·044
„ „ diluted	1·006
„ „ glacial	1·065 to 1·066
„ carbolic	1·065
„ hydrochloric	1·160
„ „ diluted	1·052
„ hydrocyanic	0·997
„ nitric	1·420
„ „ diluted	1·101
„ nitro-hydrochloric	1·074
„ phosphoric, diluted	1·080
„ sulphuric	1·843
„ „ aromatic	0·927
„ „ diluted	1·094
„ sulphurous, solution of	1·040

Alcohol, absolute	0.795
„ (rectified spirit, 84 per cent.)	0.838
„ (proof spirit, 49 per cent.)	0.920
„ amylic	0.818
Ammonia, aromatic spirit of	0.870
„ solution of	0.959
„ strong solution of	0.891
Antimony, solution of chloride of	1.470
Arsenic, hydrochloric solution of	1.009
Arsenical solution (<i>Liquor Arsenicalis</i>)	1.009
Benzol	0.850
Bismuth and ammonia, sol. of citrate of	1.122
Bromine	2.966
Chlorine, solution of	1.003
Chloroform	1.490
„ spirit of	0.871
Cinchona, liquid ext. of yellow (about)	1.100
Creasote	1.071
Ether	0.735
„ pure	0.720
Glycerine	1.250
Iron, solution of perntrate of	1.107
„ „ persulphate of	1.441
„ strong solution of perchloride of	1.440
„ tincture of perchloride of	0.992
Lead, solution of subacetate of	1.260
Lime, saccharated solution of	1.052
„ solution of chlorinated	1.035
Mercury (at 0° C. = 32° F.)	13.596
„ (at 15.55° C. = 60° F.)	13.560
„ acid solution of nitrate of	2.246
Nitre, sweet spirit of	0.845
Oil of mustard	1.015
Potash, solution of	1.058
Soda, „	1.047
„ „ ohlorinated	1.103
Squill, oxymel of	1.320
Syrup	1.330
„ of buckthorn	1.320
„ of chloral	1.218
„ of ginger	
„ of hemidesmas	1.335
„ of iodide of iron	1.385

Syrup of lemons	1.340
„ of mulberries	1.330
„ of orange-flower	1.330
„ of „ peel	
„ of phosphate of iron	
„ of poppies	1.320
„ of red poppy	1.330
„ of „ roses	1.335
„ of rhubarb	
„ of senna	1.310
„ of squill	
„ of tolu	1.330
Treacle (about)	1.400

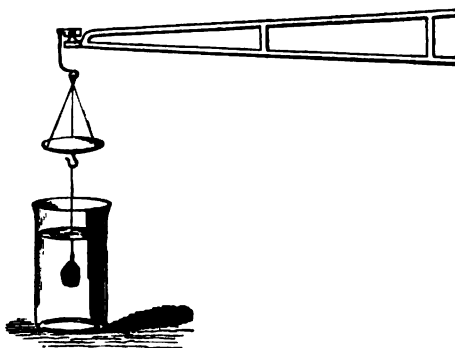
Hydrometers.—The specific gravity of liquids may be ascertained, without scales and weights, by means of an *hydrometer*—an instrument usually of glass, having a graduated stem and a bulb or bulbs at the lower part. The specific gravity of a liquid is indicated by the depth to which the hydrometer sinks in the liquid, the zero of the scale marking the depth to which it sinks in pure water. Hydrometers constructed for special purposes are known under the names of saccharometer, galactometer, elæometer, urinometer, alcoholometer. Hydrometers require a considerable quantity of liquid fairly to float them: and specific gravities observed with them are less delicate and trustworthy than those obtained by the balance, nevertheless are exceedingly useful for many practical purposes where the employment of a delicate balance would be inadmissible.

SPECIFIC GRAVITY OF SOLIDS IN MASS.

Weigh a piece (50 to 250 grains) of any solid substance heavier than water in the usual manner. Then weigh it in water, by suspending it from a shortened balance-pan by a fine thread or hair and immersing in a vessel of water (fig. 70). The buoyant properties of the water will cause the solid apparently to lose weight: *this loss in weight is the exact weight of an equal bulk of water.* The weight of the substance and the weight of an equal bulk of water being thus ascertained, a rule-of-three sum shows the proportional weight of

the substance to 1·000 of water. To express the same thing by rule, divide the weight in air by the loss of weight in water, the resulting number is the specific gravity in relation to 1 part of water, the conventional standard of comparison.

Fig. 70.



WEIGHING A SOLID IN WATER.

Verify some of the following specific gravities :—

Aluminium.	2·56
Antimony	6·71
Bismuth	9·83
Coins, English, gold	17·69
" " silver	10·30
" " bronze	8·70
Copper	8·95
Gold.	19·34
Iron	7·84
Lead.	11·36
Magnesium	1·74
Marble	2·70
Phosphorus	1·77
Platinum	21·53
Silver	10·53
Sulphur.	2·05
Tin	7·29
Zinc	7·14

Specific gravities of solid substances should be taken in water having a temperature of about 60° F. The body should be immersed about half an inch below the surface of the water; adhering air-bubbles must be carefully removed; the body must be quite insoluble in water.

(For a table of the specific gravities of a large number of fatty and resinoid substances, see the *Pharmaceutical Journal* for Oct. 11, 1879.)

SPECIFIC GRAVITY OF SOLIDS IN POWDER OR SMALL FRAGMENTS.

Weigh the particles; place them in a counterpoised specific-gravity bottle of known capacity, and fill up with water, taking care that the substance is thoroughly wetted; again weigh. From the combined weights of water and substance subtract the amount due to the substance; the residue is the weight of the water. Subtract this weight of water from the quantity which the bottle normally contains: the residue is the amount of water displaced by the substance. Having thus obtained the weights of equal bulks of water and substance, a rule-of-three sum shows the relation of the weight of the substance to 1 part of water, the specific gravity.

Or, suspend a cup, short glass tube, or bucket from a shortened balance-pan; immerse in water; counterpoise; place the weighed powder in the cup, and proceed as directed for taking the specific gravity of a solid in mass.

This operation may be conducted on fragments of any of the substances the specific gravities of which are given in the foregoing Table, or on the powdered piece of marble the specific gravity of which has been taken in mass. The specific gravity of one piece of glass, first in mass then in powder, may be ascertained; the result should be identical. The specific gravity of shot is about 11.350; sand, 2.600; mercury, 13.56.

SPECIFIC GRAVITY OF SOLIDS SOLUBLE IN WATER.

Weigh a piece of sugar or other substance soluble in water; then suspend it from a balance in the usual manner, and weigh it in turpentine, benzol, or petroleum, the specific gravity of which is known or has been previously determined;

the loss in weight is the weight of an equal bulk of the turpentine. Ascertain the weight of an equal bulk of water by calculation :—

Sp. gr. of sp. gr. of observed equal bulk
turpentine : water : ; bulk of turp. : of water.

The exact weights of equal bulks of sugar and water being obtained, the weight of a bulk of sugar corresponding to 1·000 of water is shown by a rule-of-three sum; in other words, divide the weight of sugar by that of the equal bulk of water, the quotient is the specific gravity of sugar. The stated specific gravity of sugar ranges from 1·590 to 1·607.

SPECIFIC GRAVITY OF SOLIDS LIGHTER THAN WATER.

This is obtained in a manner similar to that for solids heavier than water; but the light body is sunk by help of a piece of heavy metal, the bulk of water which the latter displaces being deducted from the bulk displaced by both; the remainder is the weight of a bulk of water equal to the bulk of the light body. For instance, a piece of wood weighing 12 grammes (or grains) is tied to a piece of metal weighing 22 grammes, the loss of weight of the metal in water having been previously found to be 3 grammes. The two, weighing 34 grammes, are now immersed, and the loss in weight found to be 26 grammes. But of this loss 3 grammes have been proved to be due to the buoyant action of the water on the lead; the remaining 23, therefore, represent the same effect on the wood; 23 and 12, therefore, represent the weights of equal bulks of water and wood. As 23 are to 12 so is 1 to ·5217. Or, shortly, as before, divide the weight in air by the weight of an equal bulk of water; ·5217 is the specific gravity of the wood. Another specimen of wood may be found to be three-fourths (·750) the weight of water, and others heavier. Cork varies from ·100 to ·300.

SPECIFIC GRAVITY OF GASES.

This operation is similar to that for liquids. A globe exhausted of air and holding from 1 to 4 litres (or quarts) is

suspended from the arm of a balance, and counterpoised by a similar flask. Gases are introduced in succession and their weights noted. A rule-of-three sum shows their specific gravity in relation to air or hydrogen, whichever be taken as a standard.

Correction of the Volume of Gases for Pressure.—The height of the barometer at the time of manipulation is noted. Remembering the fact "the bulk of a gas is inversely as the pressure to which it is subjected" (Boyle and Mariotte), a simple calculation shows the volume which the gas would occupy at 760 millimètres (or 29·922 inches), the standard pressure (30 inches is sometimes adopted as the standard in England).^{*} Thus 40 volumes of a gas at 740 millims. pressure are reduced to 39 when the pressure becomes 760 millimètres (or 90 vols. at 29 ins. barom. become 87 vols. at 30 inches).

Correction of the Volume of Gases for Temperature.—This is done in order to ascertain what volume the gas would occupy at 0° C. (32° F.) or 15·5° C. (60° F.), according to the standard taken. Gases are equally affected by equal variations in temperature (Charles). They expand about 0·3665 † per cent. ($\frac{1}{273}$) of their volume at the freezing-point of water for every C. degree (0·2036, or $\frac{1}{491}$ for every F. degree) (Regnault). Thus 8 volumes of gas at 0° C. will become 8·293 at 10° C.; for if 100 become 103·665 on being increased in temperature 10° C., 8 will become 8·293 (or if 100 become 102·036 on being increased 10° F., 8 will become 8·1629).

Vapour-density.—Vapours are those gases which condense to liquids at common temperatures. By the density of a vapour is meant its specific gravity. The density of a vapour is the ratio of any given volume to a similar volume of air or hydrogen at the same temperature and pressure. But, for convenience of comparison, this experimental specific gravity

^{*} In France the conventional standard height of the barometer is 760 millimètres at 0° C. (32° F.); in England it is 30 inches, the temperature of the mercurial column being 60° F. 760 millims. is equivalent to 29·922 inches; but the expansion of the metal between 32° F. and 60° F. increases the length of the column to 30·006 inches. The standards are therefore almost identical, difference in true length being counterbalanced by the temperature at which the length is observed.

† Corrected for the difference between the mercurial and air-thermometers the coefficient of expansion of air is 0·003656 (Miller). The coefficient of expansion of different gases varies very slightly, being somewhat higher for the more liquefiable gases.

is referred, by calculation as just described for permanent gases, to a temperature of 0°C. , and 760 millimètres barom. A teaspoonful or so of liquid is placed in a weighed flask of about the capacity of a common tumbler and having a capillary neck: the flask is heated in an oil-bath to a temperature considerably above the boiling-point of the liquid; at the moment vapour ceases to escape, the neck is sealed by a blowpipe-flame, and the temperature of the bath noted; the flask is then removed, cooled, cleaned, and weighed; the height of the barometer is also taken. The neck of the flask is next broken off beneath the surface of water (or of mercury), which rushes in and fills it, and again weighed, by which its capacity in cub. centims. is found. From these data the volume of vapour yielded by a given weight of liquid is ascertained by a few obvious calculations. The capacity of the globe having been ascertained, the weight of an equal bulk of air* is obtained by a rule-of-three sum. This weight of air is deducted from the original weight of the flask, which gives the true weight of the glass. The weight of the glass is next subtracted from the weight of the flask and contained vapour (now condensed), which gives the weight of material used in the experiment. The volume which this weight of material occupied at the time of experiment is next corrected for temperature (to 0°C.) and pressure (760 millimètres) in the manner just described. The weight of a similar volume of hydrogen is next found.† The weights of equal volumes of hydrogen and vapour being thus determined, the amount of vapour corresponding to 1 of hydrogen (the specific gravity or vapour-density) is shown by a short calculation. This process of finding the weight of a given volume of vapour is by Dumas. Gay-Lussac's consists in determining the volume of a given weight: it has been improved by Hofmann. A new

* 1 cub. centim. of air at 0°C. and 760 millims. weighs 0.001293 gramme.

† 1 litre (1000 cub. centims.) of hydrogen at 0°C. and 760 millimètres (the barometer being at 0°C.) weighs 0.0896 gramme—a volume sometimes termed a *crith* (from *κριθῆ*, *krithē*, a barley-corn—figuratively, a small weight); thus a litre of oxygen weighs 16 oriths, chlorine 35.5 criths, etc. 100 cubic inches of hydrogen at 32°F. weigh 2.265 grains; at 60°F. 2.143 grains (the barometer being 30 ins. at 60°F. in both cases). 100 cubic inches of air at 32°F. weigh 82.698 grains; at 60°F. 80.935 (barom. 30 ins. at 60°F.). 1 cubic inch of water weighs 252.5 (252.458 at 62°F. , and 30 in. bar.) grains. 1 gallon of water contains $277\frac{1}{2}$ (277.274 at 62°F.) cubic inches.

and excellent method by V. and C. Meyer consists, like that of Gay-Lussac, in determining the volume of the vapour of a given weight of a fluid or solid, but differs in the volume of the vapour being ascertained from an equal volume of air which the vapour is made to displace. (For a detailed description of this method and a drawing of the apparatus, see *Pharmaceutical Journal*, May 17, 1879.)

Experiment shows that the specific gravities of many gases and vapours on the hydrogen scale and the proportions in which they combine by weight are identical. Thus chlorine is 35.5 times as heavy as hydrogen, and 35.5 parts unite with 1 of hydrogen to form hydrochloric acid gas. Hence if the specific gravity of a gas or vapour is known, its combining-proportion may be predicated with reasonable certainty, and *vice versâ*. In applying this rule to gaseous or vaporous compounds, attention must be paid to the extent to which their constituent gases contract at the moment of combination or expand at the moment of decomposition. Thus steam is found to be composed of two volumes of hydrogen and one of oxygen, the three volumes of constituents condensing to two at the moment of combination. Hence steam may be expected to be nine times as heavy as hydrogen, which experiment confirms.

These relations may be so expressed as to include both elementary and compound gases and vapours, thus: *molecular weights and specific weights are identical*. Molecular weights represent two volumes of a gas; specific gravity conventionally represents the relative weight of a gas compared with one volume of hydrogen or air; hence the specific gravity of a gas or vapour on the H scale is found by calculation on simply dividing the molecular weight by 2; on the air-scale by dividing the hydrogen numbers by 14.44. For example:—

Name.	Molecular formula.	Molecular weight.	Specific gravity.		
			H = 2.	H = 1.	Air = 1.
Hydrogen	H ₂	2	2	1	.069
Chlorine	Cl ₂	71	71	35.5	2.460
Oxygen	O ₂	32	32	16	1.108
Nitrogen	N ₂	28	28	14	.970
Steam	H ₂ O	18	18	9	.624
Ammonia gas . . .	NH ₃	17	17	8.5	.589
Carbonic acid gas .	CO ₂	44	44	22	1.524
Alcohol (vapour) .	C ₂ H ₆ O	46	46	23	1.593
Air	...	...	28.88	14.44	1.000

These specific gravities closely correspond with those obtained by actual experiment. The specific gravity of any gas or vapour may therefore be calculated if the following data are at hand :—(a) formula, (b) atomic weight of constituent elements; these give the molecular weight, and *the molecular weight divided by 2 is the specific gravity on the hydrogen-scale*. Specific gravity on the air-scale is then deducible, if (c) the specific gravity of air (14.44) in relation to hydrogen be remembered. The absolute weight of any volume of a gas or vapour on the metric system is then obtainable if (d) the weight of a litre of hydrogen (0.0896 gramme) be known, or on the English plan by remembering (e) that 100 cubic inches of hydrogen at 60° F. weigh 2.143 grains (100 cubic inches of air at 60° F. weigh 30.935 grains).

In confirmation of these statements regarding the mutual relation of specific gravity and atomic weight, a remarkable fact may be mentioned. Regnault several years ago found the weights of 1 litre of hydrogen and oxygen to be respectively .089578 and 1.429802 gramme. The latter number divided by the former gives 15.96 as the specific gravity of oxygen. Stas, in recent experimental researches on combining-proportion, finds the atomic weight of oxygen to be not 16, but 15.96.

Exceptions to the law occur in a few compounds and in arsenicum and phosphorus, whose vapour-densities are twice that indicated by the rule. Possibly in these cases the temperature employed is insufficient to dissociate an unusually complex molecule into molecules of usual complexity. As regards compounds, and, possibly, as regards those elements in which the observed density is only half that indicated by this rule, heat may, and in some cases probably does, produce molecular dissociation (*thermolysis*) into free atoms (uniatomic molecules) or into less complex molecules.

Relation of the specific heat of elements to their atomic weights.

—Reference may here appropriately be made to a physical fact of great importance as regards molecular and atomic weights. In the earlier pages of this manual it was stated that elements do not combine chemically in haphazard proportions, but in fixed weights; and abundant evidence of the truth of the statement has already been afforded, and will also be found in this section on quantitative analysis. Secondly, it has been shown that elements do not combine in haphazard proportions by volume, but in certain constant bulks; and the

weights of these bulks have been found to be identical with the combining weights themselves. Thirdly (this is the point to which attention is now drawn), if equal amounts of heat be given to elements in the solid state (that is, to solid elements or to solid compounds of volatile elements), and the quantity of the element be increased or diminished until each is thus heated through an equal number of degrees, it will be found that the different weights of elements required are (in relation to a common standard) identical with the combining weights of the elements, and with the weights of the combining volumes of the elements. Thus, where 108 parts of silver would be employed, 207 of lead would be necessary.* Hence, in the determination of (a) combining-proportion, (b) specific gravity in gaseous state, and (c) specific heat, three distinct methods of ascertaining atomic weight are available. In cases where one method is inapplicable recourse is had to either or, if practicable, both of the others, and thus the trustworthiness of observations and generalizations placed more or less beyond question. The specific heat of a solid element is the same in the free as in the combined condition; therefore the specific heat of a molecule is the sum of the specific heats of its constituent atoms. From the specific heat of a solid compound of a volatile element (chlorine, for example) can thus be calculated the specific heat of an element in the solid state, even though the free element cannot itself be solidified. For the processes by which experimentally to determine specific heat the reader is referred to books on physics.

There is equivalency also between *electrical* and *chemical* action. The amount of electricity which would set free 127 parts of iodine would set free 80 parts of bromine.

* Obviously if equal weights of silver and lead were heated through an equal number of degrees the silver would absorb nearly twice as much heat as the lead. In fact as regards all the solid elements "specific heats and atomic weights are inversely proportional." This law was discovered by Dulong and Petit. It follows that the product of the multiplication of the figures representing the specific heat of an element with the figures representing its atomic weight is in the case of every such element the same number.

QUESTIONS AND EXERCISES.

967. Define specific weight, or as it is commonly termed, specific gravity.

968. In speaking of light and heavy bodies specifically, what standard of comparison is conventionally employed?

969. How are specific gravities expressed in figures?

970. Why should specific gravities be taken at one constant temperature?

971. How does the buoyancy of air affect the real weight of any material?

972. Describe the difference between density and specific gravity.

973. Give a direct method for the determination of the specific gravity of liquids.

974. A certain bottle holds 150 parts, by weight, of water, or 135.7 of spirit of wine; what is the specific gravity of the latter?
—*Ans.* 0.9046.

974a. An imperial fluid ounce of a liquid weighs 366½ grains; what is its specific gravity?—*Ans.* .838.

975. Equal volumes of benzol and glycerine weigh 34 and 49 parts respectively, and the specific gravity of the benzol is 0.850; what is the specific gravity of the glycerine?—*Ans.* 1.225.

976. Explain the process employed in taking the specific gravity of solid substances in mass and in powder.

977. State the method by which the specific gravity of a light body, such as cork, is obtained.

978. What modifications of the usual method are necessary in ascertaining the specific gravity of substances soluble in water?

979. How is the specific gravity of gases determined?

980. By what law can the volume of a gas, at any required pressure, be deduced from its observed volume at another pressure?

981. To what extent will 78 volumes of a gas at 29.3 inches barom. alter in bulk when the pressure, as indicated by the barometer, is 30.2 inches?

982. Write a short account of the means by which the volumes of gases are corrected for temperature.

983. At the temperature of 15° C. 40 volumes (litres, pints, ounces, cubic feet, or other quantity) of a gas are measured. To

what extent will this amount of gas contract on being cooled to the freezing-point of water (0°C)?

Answer. As 1 vol. of any gas at zero expands or contracts $\cdot 003665$ of a vol. for each rise or fall of 1°C ., 1 vol. at 0°C . if heated to 15°C . will become increased by $\cdot 054975$ (that is, $\cdot 003665$ multiplied by 15°), 1 vol. will expand to $1\cdot 054975$. Conversely, $1\cdot 054975$ vol. will contract to 1 vol. if cooled from 15°C . to 0°C . And if $1\cdot 054975$ becomes 1 in cooling through 15°C ., 40 vols. will (as found by rule of three) contract to $37\cdot 916$.

(The following five problems and solutions are from William-son's "Chemistry.")

984. 10 litres of oxygen are measured off at 14°F . Required the volume of the gas at 15°C .

Answer. The first operation must be to reduce the temperature quoted in Fahrenheit's degrees, to an equivalent value on the Centigrade scale. 14°F . is 18° below 32°F ., the freezing-point of water; and a range of 9° on the Fahrenheit scale is equal to a range of 5° on the Centigrade scale, so that the temperature at which the oxygen is measured off is -10°C . The rise of temperature up to 0° expands the gas in such proportion that its volume at 0° is to its volume at -10° as 1 is to $1 - 0\cdot 03665$, i.e. as 1 to $0\cdot 96335$. The further rise of temperature from 0°C . to 15° expands the gas in the proportion of 1 to $1 + 15 \times 0\cdot 003665$, i.e. 1 to $1\cdot 054975$. The total rise of temperature therefore expands the gas in the proportion of $0\cdot 96335$ to $1\cdot 054975$.

$$0\cdot 96335 : 1\cdot 054975 :: 10 : x;$$

$$\therefore x = \frac{10 \times 1\cdot 054975}{0\cdot 96335} = 10\cdot 95.$$

985. 230 cubic centimètres of oxygen are measured off at 14°C . and 740 millimètres mercurial pressure. Required the volume of the gas at the normal temperature and pressure (0°C . and 760 millimètres).

Answer. Let the reduction for change of temperature be made first. The proportion

$$1 + (14 \times 0\cdot 003665) : 1 :: 230 : x$$

gives

$$x = \frac{230}{1\cdot 05131} = 218\cdot 774.$$

To reduce this volume at 740 millimètres pressure to the volume corresponding to the pressure of 760 millimètres, we have the proportion

$$38 : 37 :: 218.77 : x;$$

whence

$$x = \frac{37 \times 218.77}{38} = 213.02$$

986. A litre of oxygen is confined in a glass flask at 10° C. by the atmospheric pressure, added to that of a column of mercury 60 millimètres high. The flask must be heated to 300° C. without any increase of volume taking place in the oxygen. How high must the column of mercury then be which presses on the gas, supposing the atmospheric pressure to remain constant at 760 millimètres?

Answer. The oxygen is given at 10° C. and 820 millimètres pressure. If the pressure remained constant, the rise of temperature from 10° C. to 300° C. would expand the gas in such proportion that 1.03665 volume would expand to 2.0995 volumes. In order to prevent any expansion the pressure must be increased in the same proportion, whence

$$1.03665 : 2.0995 :: 820 : x;$$

$$\therefore x = \frac{820 \times 2.0995}{1.03665} = 1660.6$$

From this total pressure the atmospheric pressure of 760 millimètres has to be deducted, leaving 900.6 millimètres as the height of the required mercurial column.

987. A litre of oxygen is required of the density of 100 at 0° C. What weight of potassic chlorate must be used for its preparation, and what total pressure must be applied to it?

Answer. The pressure required to compress oxygen from the density of 16 to that of 100 is found by the proportion

$$16 : 100 :: 760 : x;$$

$$\therefore x = \frac{76000}{16} = 4750.$$

At the pressure of 4750 millimètres of mercury the weight of a litre of oxygen (16 grammes measure 11.2 litres at 0° C. and 760 millims. pressure) is found by the proportion

$$760 : 4750 :: \frac{16}{11.2} : x;$$

whence

$$x = \frac{16 \times 4750}{11.2 \times 760} = 8.93 \text{ grammes.}$$

The weight of chlorate required for the evolution of 8.93 grammes of oxygen is found from the proportion

$$48 : 122.5 :: 8.93 : x;$$

$$\therefore x = 22.8 \text{ grammes.}$$

988. What is the volume of 12 grammes of hydrogen at 15° C.?

Answer. One gramme of hydrogen measures 11.2 litres at 0°C., therefore 12 grammes measure $12 \times 11.2 = 134.4$ litres at 0°. To find their volume at 15° C. we have the proportion

$$1 : 1 + 15 \times 0.003665 :: 134.4 : x;$$

whence

$$x = 134.4 \times 1.054975 = 141.788 \text{ litres.}$$

989. What interest for chemists have the specific heats of substances?

QUANTITATIVE ANALYSIS.

Preliminary Note.—Great care should be observed in selecting a fair sample of any bulk of material that is to be examined either by volumetric or gravimetric quantitative analysis. If the whole quantity is in separate parcels, and there is any ground for believing that the parcels differ in quality, they should, if practicable, be carefully mixed, or, technically, "bulked." Small portions should be taken from different parts of the resulting heap and well mixed in a mortar or other vessel, or, in certain cases, dissolved, and the solution well stirred or shaken. A specimen of the powder, or a portion of the solution, may then be selected for analysis.

VOLUMETRIC QUANTITATIVE ANALYSIS.

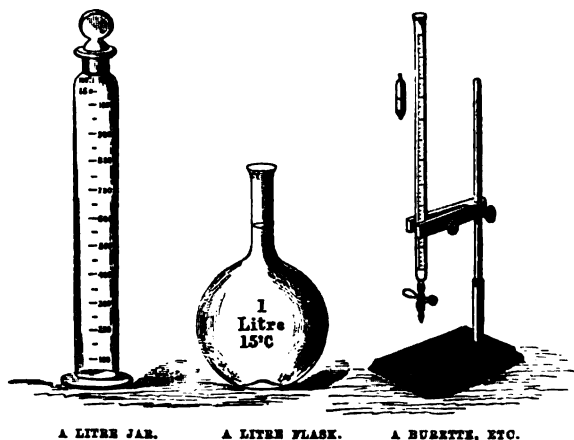
Introduction.—The operations of volumetric analysis consist (a) in carrying out some definite *chemical reaction*, already well-known to the operator, with (b) *definite quantities* of

chemicals or salts; (c) the exact termination of the reaction between the two salts or chemicals being ascertained—usually by some *chemical indicator* (litmus, starch, etc.). A portion of the chemical or salt, etc., to be tested is carefully weighed. To this is gradually added the second chemical or salt contained in the testing fluid commonly termed the Standard Volumetric Solution. The usefulness, and, indeed, the preparation of this Standard Solution is founded (as already indicated on page 609) on some accurate initial gravimetric operation. A *weighed* amount of a pure salt is dissolved in a given *volume* of water. “Accurately *measured* quantities of such a Standard Volumetric Solution will obviously contain just as definite amounts of the dissolved salt as if those amounts were actually *weighed* in a balance, and as measuring occupies less time than weighing, the volumetric operations can be conducted with great economy of time as compared with the corresponding gravimetric operations.”

Fig. 71.

Fig. 72.

Fig. 73.



A LITRE JAR.

A LITRE FLASK.

A BURETTE, ETC.

APPARATUS.

The only special vessels necessary in volumetric quantitative operations are:—1. A *litre flask* (fig. 72), which, when filled to a mark on the neck, contains at 15° C., or about 60° F., one litre (1000 cubic centimètres, *i.e.* 1000 grammes of

water*); it serves for preparing solutions in quantities of one litre. 2. A tall cylindrical *graduated litre jar* (fig. 71) divided into 100 equal parts; it serves for the measurement and admixture of decimal or centesimal parts of a litre. 3. A graduated tube or *burette* (fig. 73), which, when filled to 0, holds 100 cubic centimètres (a décilitre), and is divided into 100 equal parts; it is used for accurately measuring small volumes of liquids.

The best form of burette is Mohr's. It consists of a glass tube commonly about the width of a little finger and the length of an arm from the elbow, contracted at the lower extremity and graduated. The width and length of burettes, however, as well as the extent and fineness of their graduation, vary considerably. To the contracted portion is fitted a small piece of vulcanized caoutchouc tubing, into the other end of which a small spout made of narrow glass tube is tightly inserted. A strong wire clamp effectually prevents any liquid from passing out of the burette unless the knobs of the clamp are pressed by the finger and thumb of the operator, when a stream or drops flow at will. In place of the india-rubber tubing and clamp, a stopcock is sometimes employed; and other modes of arresting the flow of liquid may be adopted. The accurate reading of the height of a solution in the burette is a matter of great importance; it should be taken from the bottom of the curved surface of the liquid. It may be still more exactly measured by the employment of a hollow glass float or bulb (Erdmann's float—see fig. 73), of such a width that it can move freely in the tube without undue friction, and so adjusted in weight that it shall sink to more than half its length in any ordinary liquid. A fine line is scratched round the centre of the float; this line must always be regarded as marking the height of the fluid in the burette. In charging the burette, a solution is poured in, not until its surface is coincident with 0, but until the mark on the float is coincident with 0.

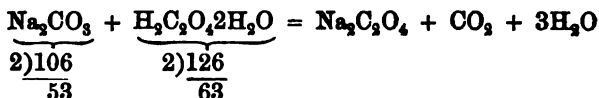
* A cubic centimètre is, strictly speaking, the volume occupied by one gramme of distilled water at its point of greatest density, namely 4° C.; metrical measurements, however, are uniformly taken at 15.55° C. (60° F.).

ESTIMATION OF ALKALIES, ETC.**VOLUMETRIC SOLUTION OF OXALIC ACID.**(Crystallized Oxalic Acid, $\text{H}_2\text{C}_2\text{O}_4 \cdot 2\text{H}_2\text{O} = 126$.)

On account of the bivalent character of the oxalic radical and the univalent character of most of the metals contained in the salts which are estimated by oxalic acid, it is convenient that each litre of the volumetric solution should contain half a molecular weight in grammes of the acid ($\text{H}_2\text{C}_2\text{O}_4 \cdot 2\text{H}_2\text{O} = 126 \div 2 = 63$).

If pure crystallized oxalic acid be at hand the solution is made by dissolving 63 grammes in water and making the volume up with more water to exactly one litre.

Pure oxalic acid, however, not being easy to obtain, the solution may be made from the commercial acid by dissolving 65 to 70 grammes in enough water to make a litre of solution and then determining the strength of this solution by a titration with pure carbonate of sodium, making use of the following memoranda:—



Pure anhydrous carbonate of sodium is easy to obtain, for commercial bicarbonate is usually of such purity that when a few grammes are heated to redness for a quarter of an hour the resulting carbonate is practically free from impurity. The bicarbonate should, however, be tested, and if more than traces of chlorides and sulphates are present these may be removed by washing a few hundred grammes, first with a saturated solution of bicarbonate of sodium, and afterwards with pure distilled water. After drying, the salt is ready for ignition.

About half a gramme of the carbonate of sodium is accurately weighed and placed in a half-pint flask, around the neck of which is tied calico or leather to protect the fingers when the heated vessel is shaken by the operator. The salt is dissolved in water to about one-third the capacity of the flask, and a few drops of the indicator, blue tincture of litmus, is added. The acid solution to be "set" or "standardized" is then poured into a burette and run therefrom into the flask until the reddened litmus indicates the presence of free acid.

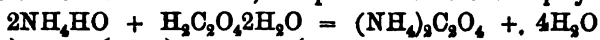
This will be due in the first place to carbonic acid liberated and remaining dissolved in the solution. The contents of the flask are therefore boiled for several minutes, when the blue colour will have returned. More acid is then run in until the mixture, after boiling, remains of a neutral colour, indicating that just enough acid has been added to complete the reaction expressed in the foregoing equation.

Let it be supposed that 6 grammes of carbonate of sodium were taken, and that this required 11 c. c. of oxalic acid solution, how many c. c. of this solution would contain 63 grammes of oxalic acid crystals; or, what is equivalent in the reaction, how many c. c. would be required to neutralize 53 grammes of carbonate of sodium? As 6 grammes Na_2CO_3 are to 11 c. c. sol., so are 53 grammes Na_2CO_3 to x c. c. sol. $x = 972$ c. c. 972 c. c. (nearly) are equivalent to 53 grammes of carbonate of sodium, and contain 63 grammes of oxalic acid.

This solution may either be used as it is or may be diluted with water, every 972 c. c. to be diluted to 1000 c. c., so that 1000 c. c. shall contain 63 grammes of oxalic acid.

The following official substances are tested by this solution according to the British and United States Pharmacopœias:—

Solutions of Ammonia.—Two or 3 grammes of dilute, or about 1 gramme of strong solution of ammonia, is a convenient quantity to operate upon. The weighing is most conveniently accomplished by taking a small stoppered bottle containing half an ounce or so of the substance, and having ascertained its total weight, transfer about the quantity desired to the flask in which the estimation is to be conducted, and again weigh the bottle with what remains in it. The difference is the exact quantity taken. The weighing of the ammonia solution having been accomplished, water is added, to about one-third the capacity of the flask (or, better, the ammonia is added to water already in the flask), and a few drops of tincture of litmus introduced. The titration is then conducted as described before, except that no heat is employed.

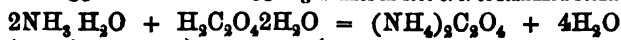


2)70

35

2)126

63 = grammes in 1000 c. c. of standard solution.



2)34

17

2)126

63 = grammes in 1000 c. c. of standard solution.

1000 c. c. of standard solution, or its equivalent of a solution of any other strength, would, according to this reaction, neutralize 17 grammes of ammonia gas (NH_3) or 35 grammes of hydrate of ammonium (NH_4HO). If 3 grammes of ammonia solution had been taken, and it had required 15 c. c. of standard oxalic acid solution, then the amount of ammonia gas or hydrate of ammonium it contained would be seen by the following calculations:—

$$\begin{array}{l} 1000 \text{ c. c.} : 17\text{NH}_3 :: 15 \text{ c. c.} : x = \cdot 255 \text{ grammes NH}_3 \\ 1000 \text{ c. c.} : 35\text{NH}_4\text{HO} :: 15 \text{ c. c.} : x = \cdot 525 \text{ grammes NH}_4\text{HO} \end{array}$$

Three grammes, then, would contain $\cdot 255$ grammes of the gas, or $\cdot 525$ grammes of hydrate of ammonium. Or in percentage:—

$$\begin{array}{l} 3 \text{ gr. sol.} : \cdot 255 \text{ gr. NH}_3 :: 100 \text{ gr. sol.} : x \text{ gr. NH}_3 = 8\cdot 5\% \text{ NH}_3 \\ 3 \text{ gr. sol.} : \cdot 525 \text{ gr. NH}_4\text{HO} :: 100 \text{ gr. sol.} : x \text{ gr. NH}_4\text{HO} = 17\cdot 5\% \text{ NH}_4\text{HO} \end{array}$$

The solution would therefore contain 8·5 per cent. of ammonia gas (NH_3) or 17·5 per cent. of hydrate of ammonium (NH_4HO). If the oxalic acid solution was not of full standard, the number of c. c. which contained 63 grammes of oxalic acid, which was, in fact, equivalent to 1000 c. c. of standard solution, would be substituted for 1000 c. c. in the preceding proportions.

A comparison should now be made with the requirements of the Pharmacopœia. It is useful to express results as percentage of substance of pharmacopœial strength in the material examined. Thus the British Pharmacopœia requires dilute ammonia solution to contain 10 per cent. of the gas (NH_3). The solution supposed to have been operated on contained 8·5 per cent. NH_3 ($10 : 8\cdot 5 :: 100 : x = 85$). Therefore it contains 85 per cent. of the dilute ammonia of the British Pharmacopœia.*

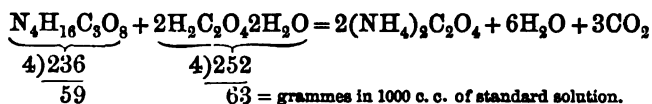
Strong Solution of Ammonia, B. P., contains 32 per cent. of ammonia gas (NH_3).

* Extremely minute quantities of ammonia—1 part in many millions of water—may be estimated volumetrically by adding excess of a colourless solution of red iodide of mercury (Nessler's test), then in a similar vessel, containing an equal amount of pure water with excess of the Nessler reagent, imitating the depth of yellow or reddish-yellow colour thus produced by adding an ammoniacal solution of known strength. The amount of ammonia thus added represents the amount in the original liquid.

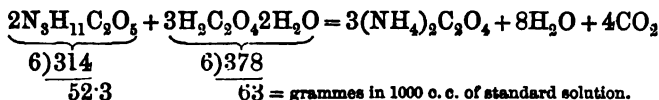
The Nessler Reagent.—A litre may be made by dissolving 80 or 40 grammes of iodide of potassium in a small quantity of hot water, adding

Note.—The calculations just described for ammonia are similar to those employed throughout volumetric analysis; they will not be repeated, therefore, in the case of every substance.

Carbonate of Ammonium.—The reactions indicated by the following equations occur between commercial carbonate of ammonium and oxalic acid :—



Or, *vide* p. 98,

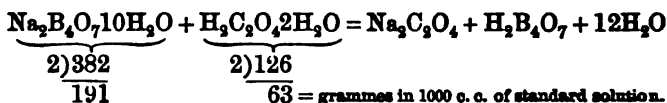


About 1 gramme is a convenient quantity to operate upon. Tincture of litmus is the indicator, and the titration is conducted at a temperature just short of boiling. The estimation is not very satisfactory, because the heat employed, while scarcely sufficient to expel the carbonic acid gas, is enough to occasion loss of ammoniacal salt. Practised analysts usually add excess of the standard acid and thus fix every trace of ammonia; then gently boil to get rid of carbonic acid gas; bring back the liquid to neutrality by an observed volume of standard alkaline solution, and deduct an equivalent volume of acid from the quantity first added. Both the British and United States Pharmacopœias require 5·9 grammes to neutralize 100 c.c. of standard solution of oxalic acid. This corresponds to 100 per cent. of carbonate having the formula $\text{N}_4\text{H}_{16}\text{C}_3\text{O}_8$.

Borax.—Two or three grammes is a convenient quantity.

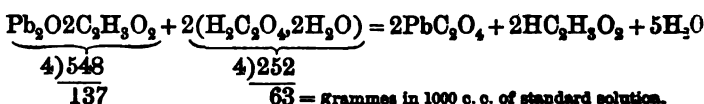
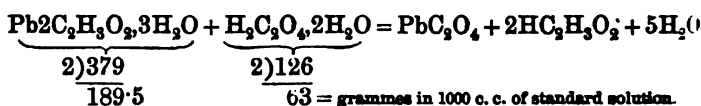
a strong hot solution of perchloride of mercury until the precipitate of mercuric iodide ceases to redissolve even by the aid of rapid stirring and heat, slightly diluting, filtering, adding a strong solution of (120 to 140 grammes of) caustic soda or (160 to 180 grammes of) caustic potash, and diluting to 1 litre. A few c. c. (5 or 6 or more) of a strong solution of perchloride of mercury are finally stirred in, the whole set aside till all precipitated red iodide has deposited, and the clear liquid decanted for use. The reaction of this Nessler test with ammonia is as follows :—





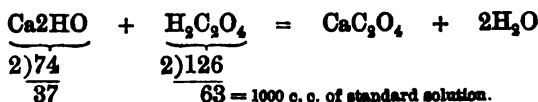
Tincture of litmus is the indicator, and the titration may be carried on without heat. The liberation of boracic acid colours the litmus wine-red. This is not regarded, the titration being continued until the bright red due to the action of free oxalic acid makes its appearance. Both the British and United States Pharmacopœias require borax to be pure (= 100 per cent.).

Lead Acetate, and Solution of Subacetate.—Operate upon about three grammes of acetate of lead, and from five to ten grammes of solution of subacetate.

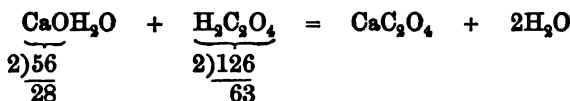


The flask in which the estimation is being conducted should contain one-third of a flaskful of water. In the case of both acetate and solution of subacetate of lead a little acetic acid should be added to prevent precipitation of basic salt on dilution. The only indicator of complete reaction is cessation of production of the precipitate—oxalate of lead. Both the British and United States Pharmacopœias require acetate of lead to be pure (100 per cent.) and solution of subacetate to contain nearly 24.9 per cent.

Lime Water, and Saccharated Solution of Lime.—Measure about half a litre of lime water for the estimation, and of saccharated solution weigh about 25 grammes. The following equations, etc., are quantitative expressions of the reactions:—

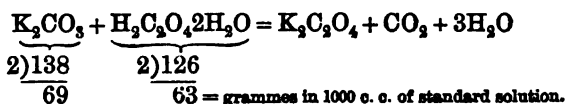
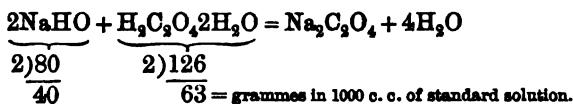
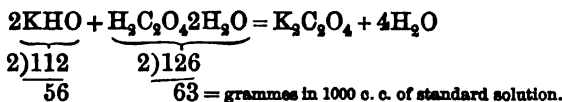


Or,

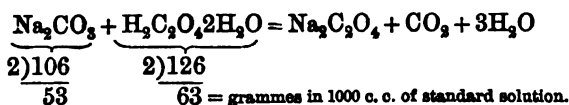
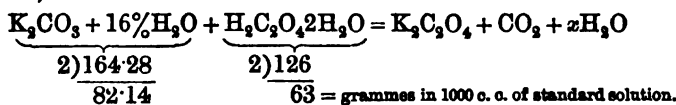


Litmus is used as an indicator.

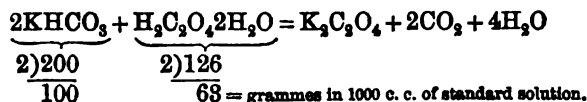
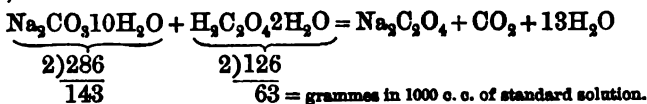
Caustic Potash and Soda. Potassium and Sodium Carbonates and Bicarbonates.—Litmus is the indicator throughout, and heat is used in all cases, for the caustic alkalies always contain some carbonate.

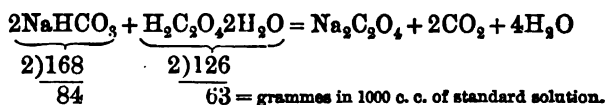


Or,



Or,



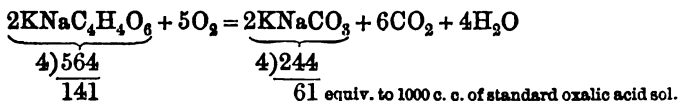
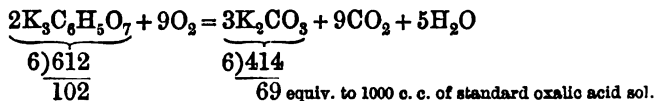
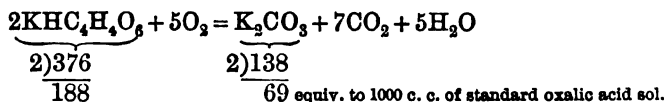
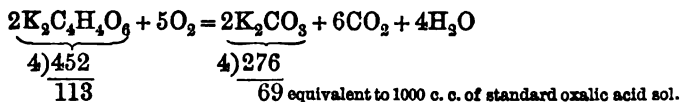


Convenient quantities to operate with are: Of caustic potash, 1 gramme; caustic soda, .5 to 1 gramme; potassium carbonate, or bicarbonate, 1 to 2 grammes; sodium carbonate, or bicarbonate, 2 to 3 grammes; dried sodium carbonate, .5 to 1 gramme; and of solutions a corresponding quantity. The British and United States Pharmacopœial requirements are: Caustic potash or soda, 90 to 100 per cent. of KHO or NaHO ; potassium carbonate, 98 to 100 per cent. of $\text{K}_2\text{CO}_3 + 16$ per cent. H_2O ; sodium carbonate, 96 to 100 per cent. of $\text{Na}_2\text{CO}_3 \cdot 10\text{H}_2\text{O}$; potassium and sodium bicarbonates, 100 per cent., respectively, of KHCO_3 and NaHCO_3 . The requirements for solutions of these substances are not exactly the same in Great Britain as in the United States.

The strength of soda ash is often reported in terms of "soda"—that is, oxide of sodium ($\text{Na}_2\text{O} = 62$). The old molecular weight of carbonate of sodium, 54 (it should have been 53), derived from that of "soda," 32 (it should have been 31), is still employed in Great Britain in reporting the strength of soda-ash. The true amount of soda equivalent to 54 parts of carbonate is 31.41 parts. A modern analyst having found the true amount of soda in a sample of soda-ash is expected by some manufacturers to report 31 as 31.41 parts, or 53 of carbonate as 54, and other quantities in proportion to these figures.

Tartrates and Citrates of Potassium and Sodium.—When tartrates or citrates of alkali-metals are burned in the open air the whole of the metal remains in the form of carbonate. Each molecular weight of a neutral tartrate gives one molecular weight of carbonate, and every two molecular weights of an acid tartrate give one molecular weight of carbonate. Advantage is taken of these reactions to estimate indirectly the quantity of citrate or tartrate in presence of substances with which they are generally associated. One to two grammes of any of these salts is a convenient quantity to operate upon. The ignition may be conducted in a platinum or porcelain crucible. A low red heat only should be used, and the vessel removed when complete carbonization has been effected—that is to say when nothing remains but the carbonate and free

carbon. The mixture is in this case treated with hot water, and the carbon separated by filtration. If too little heat has been used, and carbonization is not complete, the filtrate will be more or less coloured. If this should be the case the operation must be repeated with a fresh quantity of material. The carbonate is titrated in the usual way. The following equations, etc., explain the reactions:—



It will be readily understood that in the first (for example) of the reactions just expressed, 113 weights of tartrate of potassium are equivalent to 69 weights of carbonate of potassium; and as in a previous reaction it has been shown that 69 weights of carbonate of potassium are equivalent to 63 weights of oxalic acid, it follows that 113 weights of tartrate of potassium are equivalent to 63 weights of oxalic acid. Let these weights be grammes, and then 113 grammes of tartrate of potassium are equivalent to 63 grammes of oxalic acid, or to 1000 c. c. of the standard solution of oxalic acid. If the substance estimated be a crude sample of tartrate of potassium, and the number of c. c. of oxalic acid used has been 15 c. c., then as 1000 c. c. of the acid solution are to 113 grammes of tartrate of potassium, so are 15 c. c. of the solution to 1.695 grammes of tartrate of potassium. Now if the weight of the sample taken was 2 grammes, then as 2 grammes of the sample contain 1.695 of real tartrate of potassium, 100 will

contain $x = 84.75$ per cent. of real tartrate. These salts are required to be 100 per cent. pure by both the British and United States Pharmacopœias. Trade samples are practically pure as a rule.

Notes.

Alkalimetry.—The foregoing processes are often spoken of as those of *alkalimetry* (the measurement of alkalis).

Neutral solution of litmus is prepared by digesting the commercial fragments in about fifteen or twenty times their weight of water for a few hours, decanting, dividing into two equal portions, adding acid to one till it is faintly red, then pouring in the other and mixing. The solution may be kept in a stoppered bottle, and occasionally exposed to the air. It should never be filtered, but gradually allowed to deposit.

Standard sulphuric acid may be used in the place of oxalic acid, 1000 c. c. of the liquid containing half of the molecular weight of the pure acid in grammes. It is prepared by diluting oil of vitriol with from three to four times its bulk of distilled water, ascertaining how much of the acid liquid is required to exactly neutralize $\frac{1}{10}$ of the molecular weight of pure carbonate of sodium, taken in grammes (5.3), and adding water until the observed volume of acid is increased to 100 c. c., the whole of the fluid being similarly diluted.

Weighing.—In the case of substances which are liable to alter by exposure to air, it is important that a selected quantity should be quickly weighed, rather than selected weights be accurately balanced by material, the former operation occupying much the shorter time.

Salts other than the official may be quantitatively analysed by the volumetric solutions of the Pharmacopœia, slight modifications of manipulation even enabling the processes to be adapted to fresh classes of salts. Ample instructions for extending operations in this manner will be found in Sutton's "Handbook of Volumetric Analysis."

QUESTIONS AND EXERCISES.

990. Describe the various pieces of apparatus used in volumetric determinations.

991. One hundred cubic centimètres of solution of oxalic acid contain 6.3 grammes of the crystallized salt; work sums showing what weights of bicarbonate of potassium and anhydrous carbonate of sodium that volume will saturate.—*Ans.* 10 grammes and 5.3 grammes.

992. Show what weight of hydrate of potassium is contained in solution of potash 48.02 grammes of which are saturated by 50 c. c. of the standard solution of oxalic acid.—*Ans.* 5.83 per cent.

993. Calculate the percentage of hydrate of calcium in lime-water 438 grammes of which are neutralized by 20 c. c. of the volumetric solution of oxalic acid.—*Ans.* 0.1689.

994. Eight grammes of a sample of Rochelle salt, after ignition, etc., require 54.3 c. c. of the official oxalic acid solution for complete saturation; work sums showing what is the centesimal proportion of real salt present.—*Ans.* 95.7.

ESTIMATION OF ACIDS.

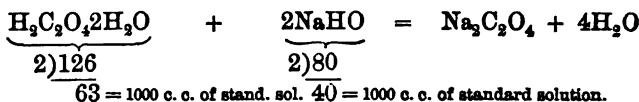
In the previous experiments a known amount of an acid has been used in determining unknown amounts of alkalies. In those about to be described a known amount of an alkali is employed in estimating unknown amounts of acids. The alkaline salt selected may be either a hydrate or a carbonate, but the former is to be preferred; for the carbonic acid set free when a strong acid is added to a carbonate, interferes to some extent with the indications of alkalinity, acidity, or neutrality afforded by litmus. The alkali most convenient for use is soda, a solution of which has probably already been made the subject of experiment in operations with the standard solution of oxalic acid. It should be kept in a stoppered bottle and exposed to air as little as possible.

VOLUMETRIC SOLUTION OF SODA.

(Hydrate of Sodium, $\text{NaHO} = 40$.)

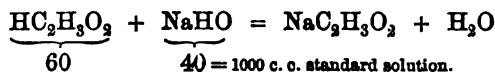
This aqueous solution of soda is most conveniently made of such a strength that each 1000 c. c. contains one molecular weight in grammes of the alkali ($\text{NaHO} = 40$). It will be seen from the following equation that 40 grammes of soda convert 63 grammes of oxalic acid into neutral oxalate of sodium. Therefore one litre of this solution, containing 40 grammes

of soda, will form a neutral solution of oxalate with one litre of standard oxalic acid solution, or with a chemically equivalent quantity of oxalic acid solution of any other strength:—



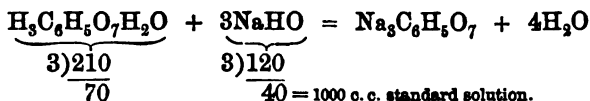
If pure soda were at hand, it would only be necessary to weigh 40 grammes, dissolve this in water, and dilute to one litre. But pure soda cannot readily be produced. Therefore weigh about 45 grammes of hydrate of sodium of trade, and add water to one litre. When dissolved take, say, 15 c. c., dilute with more water in a flask, add a few drops of tincture of litmus and titrate with oxalic acid solution of known strength. Suppose that the volume of standard acid solution required to neutralize the 15 c. c. of soda solution the strength of which is to be estimated has been 14 c. c., or an equivalent amount of acid solution of another strength; then, how many c. c. of soda solution is equivalent to 1000 c. c. of standard acid solution; or, what comes to the same thing, how many c. c. of soda solution contains 40 grammes of real soda (NaHO)? As 15 c. c. stand. acid are to 14 c. c. soda sol., so are 1000 c. c. stand. acid to x c. c. $x = 933$ c. c. 933 c. c. of the soda solution contain, therefore, 40 grammes of soda. This may either be diluted, every 933 c. c. to 1000 c. c., so that it may be standard (1000 c. c. = 40 grammes NaHO), or the solution may be used without dilution (933 c. c. = 40 grammes NaHO). It has already been mentioned that soda nearly always contains carbonate. To remove resulting carbonic acid, therefore, gentle heat should be employed towards the close of each titration in all the estimations with this solution. Litmus is used throughout as an indicator of completion of the reaction. The following substances are officially estimated with this solution. The list admits of considerable extension (*see* Sutton's "Volumetric Analysis").

Acetic Acid.—Operate upon about 1 gramme of glacial acid, about 20 grammes of dilute acid, or about 3 grammes of ordinary acetic acid.



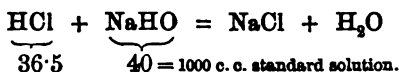
Acetic Acid, B. P., should contain 33 per cent. of real acid ($\text{HC}_2\text{H}_3\text{O}_2$). Dilute Acetic Acid, B. P. and U. S. P., 4·27 per cent. Glacial Acetic Acid, B. P., 98·8 per cent.

Citric Acid.—Operate on about 1 gramme. The reaction is expressed by the following equation, etc. :—



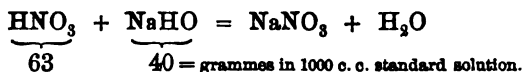
Citric Acid, B. P. and U. S. P., should be pure (= 100 per cent. $\text{H}_3\text{C}_6\text{H}_5\text{O}_7\text{H}_2\text{O}$).

Hydrochloric Acid.—Operate on from 1 to 2 grammes of the concentrated acid, or on about 4 grammes of the dilute acid.



Hydrochloric Acid, B. P. and U. S. P., should contain 31·8 per cent. of real acid (HCl), and Dilute Hydrochloric Acid, B. P., 10·58 per cent.

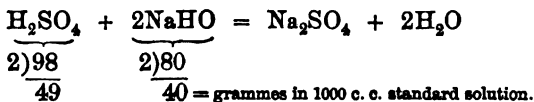
Nitric Acid.—Operate on from 1 to 2 grammes of concentrated, or on from 4 to 5 grammes of dilute acid.



Nitric Acid, B. P. and U. S. P., should contain 70 per cent., and Dilute Nitric Acid, B. P., 12·81 per cent. of real acid (HNO_3).

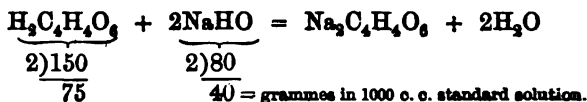
Nitrohydrochloric Acid.—Operate upon 4 to 5 grammes of dilute acid. The reaction is a complex one. The British Pharmacopoeia requires that 383 grammes of the dilute acid should neutralize 1000 c. c. of standard soda solution.

Sulphuric Acid.—Operate upon from 5 to 1 gramme of concentrated acid, or from 4 to 5 grammes of either Dilute or Aromatic Sulphuric Acid.



Sulphuric Acid, B. P. and U. S. P., should contain 96·8 per cent.; Dilute, B. P., 13·65 per cent.; and Aromatic, 13·36 per cent. of real acid (H_2SO_4).

Tartaric Acid.—Operate upon about 1 gramme of the acid. The following equation, etc., represents the reaction:—



Tartaric Acid, B. P. and U. S. P., should contain 100 per cent. of $\text{H}_2\text{C}_4\text{H}_4\text{O}_6$.

Notes.—1. Pure acetates, citrates, tartrates, and some other organic salts have an alkaline action on litmus, but not to an important extent. If the soda solution be added to acetic, citric, or tartaric acid, containing litmus, until the liquid is fairly blue, the operator will obtain trustworthy results. In delicate experiments turmeric, "eosin," "phenolphthalein," etc., may be used instead of litmus.

2. The operations for the quantitative analysis or measurement of acids are often collectively spoken of under the name of *acidimetry*.

QUESTIONS AND EXERCISES.

995. Calculate the percentage of real acid present in diluted sulphuric acid 30 grammes of which are neutralized by 84 c. c. of the official volumetric solution of soda.—*Ans.* 13·72.

996. Show how much real nitric acid is contained in a solution 36 grammes of which are saturated by 94 c. c. of the standard solution of soda.—*Ans.* 16·45 per cent.

ESTIMATION OF ACIDULOUS RADICALS PRECIPITATED BY NITRATE OF SILVER.

The purity of many salts and the strength of their solutions may be determined by this process; but at present only three official substances (namely, diluted hydrocyanic acid, bromide of potassium, and arseniate of sodium) are quantitatively analysed by standard solution of nitrate of silver.

STANDARD SOLUTION OF NITRATE OF SILVER.

(Nitrate of Silver, $\text{AgNO}_3 = 170$.)

Dissolve 17 grammes of crystals of pure nitrate of silver in one litre of water. 1000 c. c. of this solution contain $\frac{1}{10}$ of the molecular weight in grammes of nitrate of silver. It is therefore a decinormal solution.

Pure crystals of nitrate of silver can readily be obtained. When this is not the case and pure chloride of sodium is at hand, a solution may be made of approximate strength and then be standardized by means of that salt. The method may be thus indicated:—

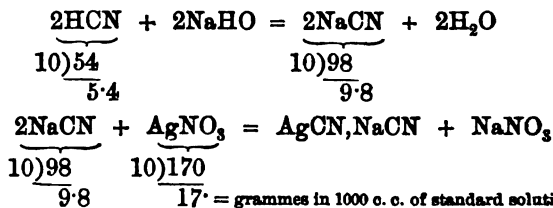
$$\begin{array}{rcl} \text{NaCl} & + & \text{AgNO}_3 = \text{AgCl} + \text{NaNO}_3 \\ 10)58.5 & 10)170 & \\ 5.85 & 17 & \end{array}$$

17 = grammes in 1000 c. c. of standard solution.

Take rather less than .1 gramme of the chloride of sodium (NaCl), and dissolve in water. The salt (AgCl) precipitated in the reaction is an insoluble salt, and the end of its precipitation will serve as a good indication of the completion of the reaction. A better indicator, however, is a few drops of neutral chromate of potassium (which should previously be purified by recrystallization). The nitrate of silver does not act upon the chromate until all the chloride is converted into chloride of silver, after which a deep red precipitate of chromate of silver is produced. This indication is extremely delicate, and in practice is noticed when the white colour due to chloride of silver changes to yellowish from formation of the first traces of chromate of silver. The titration being accomplished, suppose that .1 gramme of the chloride of sodium has taken 17 c. c. of the nitrate of silver solution of unknown strength; how many c. c. of the solution are equivalent to 5.85 of the chloride of sodium; that is, how many c. c. of solution contain 17 grammes of nitrate of silver? As .1 gramme of NaCl is to 17 c. c., so are 5.85 NaCl to x c. c. = 994 c. c. 994 c. c. of the solution of nitrate of silver are equivalent, therefore, to 1000 c. c. of official standard solution, and contain 17 grammes of the nitrate of silver. They may be diluted to 1000 c. c. if desired.

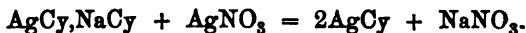
Hydrocyanic Acid.—Three to four grammes of the dilute acid form a convenient quantity to operate upon. The HCN

is first converted into KCN or NaCN, with potash or soda. The following equations, etc., explain the reactions :—



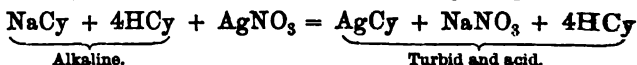
It is seen that 5.4 grammes of real hydrocyanic acid (HCy) are equivalent to 9.8 grammes of cyanide of sodium, and represent 17 grammes of nitrate of silver, or 1000 c. c. of standard solution of nitrate of silver.

The cyanide of sodium having been obtained, the titration is carried on until it is converted into the soluble double salt (NaCy, AgCy), immediately after which a permanent turbidity occurs, due to precipitation of cyanide of silver, thus :—



This turbidity affords a delicate and satisfactory proof of the completion of the above reaction.

There is, however, a difficulty in the conversion of the acid into the cyanide (Siebold), to which it is necessary to pay particular attention. Tincture of litmus is added to the acid diluted largely with water, and the soda poured in. Owing to the strong alkaline reaction of the cyanide of sodium formed, the mixture becomes blue when only a small proportion of the acid has been converted. If then the titration be conducted until the turbidity appears, only the cyanide of sodium will be estimated, leaving free hydrocyanic acid still unacted upon. Indeed, cyanide of sodium may be estimated in presence of hydrocyanic acid in this way. Thus the following reaction (expressed approximately) might occur :—

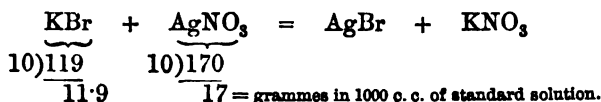


In this case only one-fifth of the acid originally present would be estimated. The mixture would, however, become acid. If this acidity be prevented all difficulty is overcome. The following details (Senier) will be found to answer well. To the diluted hydrocyanic acid add soda solution to a strong

alkaline reaction, determined by means of tincture of litmus. Then add the silver solution drop by drop from the burette, when in most cases the mixture will become acid. When it does so, add more soda solution, and repeat this process until the final reading, when the solution must be alkaline. In this way the addition of too much soda at the commencement, which would use up silver solution and make the reading a trifle too high, is avoided.

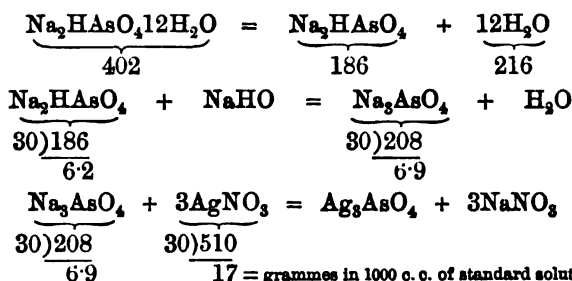
Dilute Hydrocyanic Acid, B. P. and U. S. P., should contain 2 per cent. of real acid (HCN).

Bromide of Potassium.—Operate upon rather less than .1 gramme, and conduct the titration in the same manner as with chloride of sodium, using chromate of potassium as an indicator of the close of the reaction.



Both the British and United States Pharmacopœias require bromide of potassium to contain 100 per cent. of the salt.

Arseniate of Sodium.—The crystals of $\text{Na}_2\text{HAsO}_4 \cdot 12\text{H}_2\text{O}$ lose water so readily that they cannot be relied upon to be represented by that formula. The amount of water of crystallization is estimated by drying at 300°F . In analysing, about enough soda should be added to convert the acid arseniate into neutral arseniate and the titration be then continued to the point at which a precipitate ceases to form. The reactions are shown in the following equations:—



It will be evident from the above that 402 grammes of the hydrous arseniate ($\text{Na}_2\text{HAsO}_4 \cdot 12\text{H}_2\text{O}$) are equivalent to 186

grammes of the anhydrous (Na_2HAsO_4); in other words, that crystals of the former lose 53.73 per cent. of water by desiccation. It is evident also that 6.2 grammes of anhydrous arseniate are equivalent to 17 grammes of nitrate of silver, or 1000 c. c. of standard solution of nitrate of silver. In practice about 12 grammes of the anhydrous salt are taken, dissolved in the usual quantity of water, and about .7 c. c. of standard solution of soda is added. The titration is then conducted until the precipitation is completed. Officially, both the crystallized and anhydrous salt (B. P.) are to be absolutely pure. (Another method of estimating arseniate of sodium is to treat it with sulphurous acid, and, after boiling—to expel excess of the latter—to titrate the resulting arsenite with volumetric solution of iodine.)

Spirit of Wine (*Spiritus Rectificatus*, B. P.) may contain traces of amylic alcohol and aldehyd; these may be detected by nitrate of silver, which is reduced by them to the metallic state. Any quantity beyond a mere trace of such bodies renders spirit of wine too impure for use in medicine. "Four fluid ounces with thirty grain-measures (about two c. c.) of the volumetric solution of nitrate of silver exposed for twenty-four hours to bright light, and then decanted from the black powder which has formed, undergoes no further change when again exposed to light with more of the test."

QUESTIONS AND EXERCISES.

997. Explain the volumetric method of estimating the strength of aqueous solutions of hydrocyanic acid.

998. Work a sum showing how much nitrate of silver will indicate, by the official volumetric process, the presence of 1 part of real hydrocyanic acid. *Ans.* 3.148 parts.

ESTIMATION OF SUBSTANCES READILY OXIDIZED.

Any deoxidizer, that is, any substance which quickly absorbs a definite amount of oxygen or is susceptible of any equivalent action, may be quantitatively tested by ascertaining how much of an oxidizing agent of known power must be added to a given quantity before complete oxidation is effected. The oxidizing agents employed for this purpose in the British Pharmacopœia are iodine and the red chromate of potassium.

Permanganate of potassium is often used for the same purpose. Iodine acts indirectly, by taking hydrogen from water and liberating oxygen; the red chromate of potassium directly, by the facility with which it yields three-sevenths of its oxygen—as indicated by the equations and statements given on p. 666; permanganate of potassium by affording five-eighths of its oxygen in presence of acid,



STANDARD SOLUTION OF IODINE.

(Iodine, I = 127.)

If pure iodine be not at hand, it may be prepared by mixing the commercial article with about a fourth of its weight of iodide of potassium and subliming. Sublimation may be effected by gently warming the mixture in a beaker, the mouth of which is closed by a funnel; the iodine vapour condenses on the funnel; while fixed impurities are left behind, and any chlorine which the iodine may contain is absorbed by the iodide of potassium, an equivalent quantity of iodine being liberated. Small quantities may be similarly treated between two watch-glasses, placed edge to edge. Any trace of moisture in the resublimed iodine is removed by exposure for a few hours under a glass shade near a vessel containing oil of vitriol.

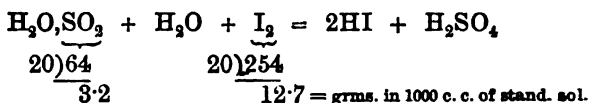
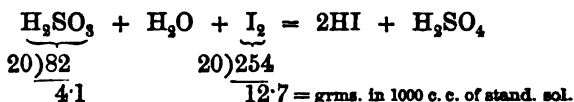
Place 12·7 grammes of pure iodine and about 18 grammes of pure iodide of potassium (an aqueous solution of which is the best solvent of iodine; the salt plays no other part in these operations) in a litre flask, add a small quantity of water, and agitate until the iodine is dissolved; dilute to 1 litre.

The following substances are officially estimated by this volumetric solution:—

Sulphurous Acid.—Operate on about ·5 of a gramme of the acid, and dilute with water as usual. If the sulphurous acid be diluted to a less degree than ·04 or ·05 per cent., there will be some risk of the sulphuric acid formed being again reduced to sulphurous acid, with liberation of iodine. In delicate experiments the distilled water used for dilution should previously be freed from air by boiling, to prevent the small amount of oxidizing action which dissolved air would exert. The solution of iodine is then added until a slight permanent brown tint is produced, showing the presence of free iodine.

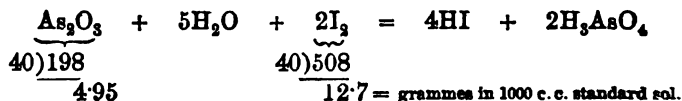
A better indicator of the termination of the reaction is mucilage of starch, which gives a blue colour with the slightest trace of free iodine.

The following equations, etc., show the reaction that takes place :—



The official (B. P.) sulphurous acid should contain 9·2 per cent. of sulphurous anhydride (SO_2).

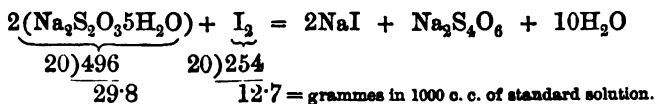
Arsenic.—About 1 gramme of *solid arsenic*, accurately weighed, should be dissolved in the usual quantity of water, heated to boiling, by help of about 5 gramme of bicarbonate of sodium. The arsenious acid is only partly, if at all, converted into arsenite or arseniate of sodium; but the iodine reaction occurs more readily in an alkaline solution. When the liquid is quite cold, mucilage of starch is added, and the iodine solution allowed to flow in until, after well stirring, a permanent blue colour is produced.—The official *arsenical solution* already containing some carbonate of sodium, requires somewhat less bicarbonate. 10 grammes is a convenient quantity to operate upon. To this should be added the usual quantity of water and about 3 of a gramme of bicarbonate of sodium. After boiling and cooling the titration is carried on as before.—About 10 grammes of the official *solution of arsenic in dilute hydrochloric acid*, is also a convenient quantity to operate upon. This quantity requires about 6 of a gramme of bicarbonate of sodium. The usual quantity of water is added, and the titration performed as before. The following equation exhibits the reaction :—



Arsenic, B. P. and U. S. P., should contain 100 per cent., and both Solutions, B. P., contain 9 per cent. of arsenic.

Hyposulphite of Sodium.—About 4 of a gramme is a convenient quantity to employ. It is dissolved in water, starch mucilage added, and the iodine solution slowly run in, the whole being frequently stirred, until a permanent blue colour is produced.

In the previous reactions iodine has acted as an indirect oxidizing agent by uniting with the hydrogen and thus liberating the oxygen of water. In the present case it unites with an analogue of hydrogen, namely sodium, a new salt (tetrathionate of sodium) being simultaneously produced, thus:—



The British Pharmacopœia requires absolute purity in the case of hyposulphite of sodium.

QUESTIONS AND EXERCISES.

999. Give equations illustrative of the reactions on which the use of a standard volumetric solution of iodine is based.

1000. From what point of view may iodine be regarded as an oxidizing agent?

1001. What reagent indicates the termination of the reaction between deoxidizing substances and moist iodine?

1002. How much sulphurous acid gas will cause the absorption of 2.54 parts of iodine in the volumetric reaction?—*Ans.* 64.

1003. What quantity of iodine will be required, under appropriate conditions, to oxidize 5 parts of arsenic?—*Ans.* 12.828.

1004. Find by calculation the amount of hyposulphite of sodium which will react with 13 parts of iodine in volumetric analysis.—*Ans.* 25.389.

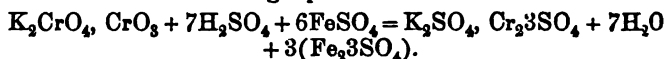
VOLUMETRIC SOLUTION OF RED CHROMATE OF POTASSIUM.

(Red Chromate of Potassium, $\text{K}_2\text{Cr}_2\text{O}_7 = 295$.)

One molecule of red chromate of potassium in presence of an acid, under favourable circumstances, yields four atoms of oxygen to the hydrogen of the acid, leaving three available

either for direct oxidation or for combination with the hydrogen of more acid, an equivalent proportion of acidulous radical being liberated for any required purpose.

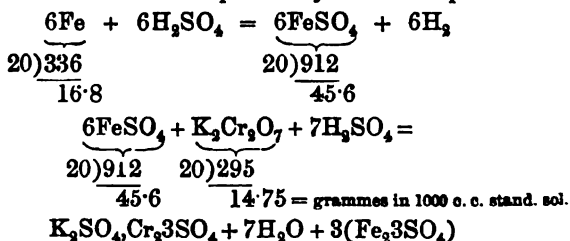
When used as a volumetric agent, the red chromate always yields the whole of its oxygen to the hydrogen of the accompanying acid, a corresponding quantity of acidulous radicals being set free—four-sevenths of this radical immediately combining with the potassium and chromium of the red chromate, three-sevenths becoming available. Ferrous may thus be converted into ferric salts with sufficient rapidity and exactitude to admit of the estimation of an unknown quantity of iron by a known quantity of the red chromate. As one atom of any liberated bivalent acidulous radical will convert two molecules of ferrous into one of ferric salt, one molecule of red chromate causes six of ferrous to become three of ferric, as shown in the following equation:—



The volumetric solution is made by dissolving 14.75 grammes ($\frac{1}{10}$ of a molecular weight in grammes) of red chromate of potassium in water, and diluting to one litre. It is used in determining the strength of the ferrous preparations. It is known that the whole of the ferrous has been converted to ferric salt when a small drop of the liquid placed in contact with a drop of a very dilute solution of ferridcyanide of potassium, on a white plate, ceases to strike a blue colour.

If the red chromate employed in making this standard solution is not known to be pure and dry, the strength of the solution may be checked by dissolving a small accurately weighed piece of pianoforte wire (0.4 or 0.5) in diluted sulphuric acid in a small flask, warming, and then running in the solution of red chromate until conversion is effected.

The reactions which take place may be thus expressed:—

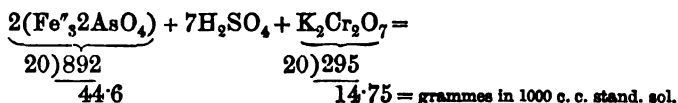


It is evident that 16·8 grammes of iron are equivalent in the reactions to 14·75 of red chromate or 1000 c. c. of standard solution of the chromate. Now suppose that 0·5 of a gramme of pianoforte wire has been employed, and the quantity of solution of red chromate of unknown strength used has been 28 c. c. How many c. c. of this solution contains 14·75 of red chromate, that is, how many c. c. must be required to oxidize ferrous salt containing 16·8 of iron? As ·5 of iron is to 28 c. c. sol., so are 16·8 of iron to x c. c. sol. = 941 c. c. Of the supposed solution, then, 941 c. c. would contain 14·75 grammes of red chromate, and would be equivalent to 1000 c. c. of standard solution. It might be employed without being diluted, or, better, be diluted to official standard strength.

Special care should be taken in all these estimations of substances readily oxidized to avoid atmospheric oxidation. Flasks may usually be loosely corked, or corked closely with a gas exit tube passing just beneath a little mercury, and in all cases the estimation should be performed quickly. When standardizing with iron wire any slight oxidation may be remedied by a fragment of zinc, the last portions of which must be removed or dissolved before the titration is commenced.

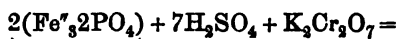
The ferrous salt in the following substances is estimated officially by this solution.

Arseniate of Iron.—Operate upon 1 to 2 grammes. Dissolve in excess of dilute sulphuric or hydrochloric acid. Sulphuric acid is preferable in most cases, because ferrous sulphate absorbs oxygen much less readily than ferrous chloride. The reaction that occurs is shown in the following equation, the ferrous arseniate being converted into ferric arseniate :—



Arseniate of iron, B. P., is supposed to contain 37·9 per cent. of ferrous arseniate. The compound is more nearly a ferric than a ferrous arseniate.

Phosphate of Iron.—Operate upon 1 to 2 grammes. Proceed as with arseniate. The following equation indicates the reaction, the ferrous phosphate being converted into ferric phosphate :—



20)716

20)295

35·8

14·75 = 1000 c. c. of standard sol.



The official (B. P.) requirement is nearly 45 per cent. of real ferrous phosphate.

Saccharated Carbonate.—Proceed as with arseniate, using about the same quantity :—



20)696

20)295

34·8

14·75 = 1000 c. c. of standard sol.



The official (B. P.) strength is 37 per cent. Trade samples yield from 20 to 30, and sometimes 35 per cent., according to the care with which oxidation has been prevented. The theoretical percentage obtainable from the ingredients is 45·5, the quantity that would be present if the compounds were anhydrous and unoxidized, conditions never obtained in practice. Howie has suggested that as hydrochloric acid is known to so rapidly convert ordinary sugar into inverted sugar as to render it easily attacked by chromic acid, while phosphoric acid very slowly affects sugar, the latter acid instead of the former should be employed in dissolving the saccharated carbonate of iron for volumetric analysis. Another mode of eliminating the action of sugar is to char with oil of vitriol before analysing.

Magnetic Oxide of Iron.—Use about the same quantity, and proceed as with arseniate or phosphate. The reaction may thus be shown :—

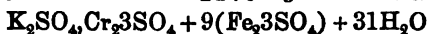


20)1392

20)295

69·6

14·75 = grms. in 1000 c. c. of standard sol.



Or,

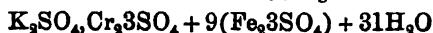


20)432

20)295

21·6

14·75 = grms. in 1000 c. c. of stand. sol.



Absolutely pure magnetic oxide of iron contains 31 per cent. of ferrous oxide. Oxidation occurs, however, during manufacture, as in the case of the ferrous salts just described. The British Pharmacopœia recognises magnetic oxide containing nearly 25 per cent. of ferrous oxide.

Note.—The use of this volumetric solution in quantitative analysis admits of great extension. The student should at least employ it in the case of a few iron ores.

QUESTIONS AND EXERCISES.

1005. Write equations explanatory of the oxidizing-power of red chromate of potassium.

1006. One hundred cubic centimètres of an aqueous solution of red chromate of potassium contain $\frac{1}{30}$ of the molecular weight of the salt in grammes; with what weight of metallic iron, dissolved in hydrochloric acid, will this volume react?—*Ans.* 1.68 gramme.

1007. If 8.34 grammes of impure crystallized ferrous sulphate, dissolved in acidulated water, require 93 c. c. of the standard solution of chromate for complete conversion into ferric salt, what percentage of ferrous sulphate is present?—*Ans.* 93.

1008. Work a sum showing how much red chromate of potassium is required for the conversion of 10 parts of ferrous sulphate into ferric salt.—*Ans.* 1.768.

1009. Show what quantity of pure ferrous carbonate is indicated by 1.475 part of red chromate as applied in volumetric analysis.—*Ans.* 3.48.

1010. Prove what amount of official saccharated carbonate of iron is equivalent to .7375 part of red chromate in the volumetric reaction.—*Ans.* 4.7.

ESTIMATION OF SUBSTANCES READILY DEOXIDIZED.

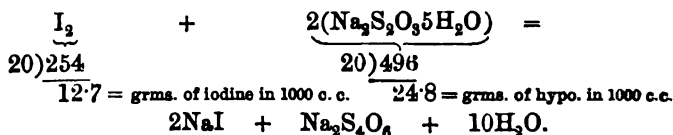
Any substance which quickly yields a definite amount of oxygen may be quantitatively tested by ascertaining how much of a deoxidizing agent of known power must be added to a given quantity before complete deoxidation is effected. The chief compounds which may be used for this absorption of oxygen (deoxidizers or reducing agents, as they are commonly termed) are hyposulphite of sodium, sulphurous acid, ferrous

sulphate,* oxalic acid, arsenious acid. The first-named is officially employed; it is only used in the estimation of free iodine, and, indirectly, of chlorine and chlorinated compounds. Iodine and chlorine are regarded as oxidizing agents, because their great affinity for hydrogen enables them to become powerful indirect oxidizers in presence of water.

STANDARD SOLUTION OF HYPOSULPHITE OF SODIUM.

(Crystallized Hyposulphite of Sodium, $\text{Na}_2\text{S}_2\text{O}_3 \cdot 5\text{H}_2\text{O} = 248$.)

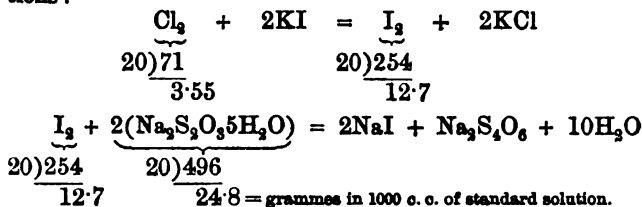
Dissolve about 27 grammes of hyposulphite of sodium in a litre or less of water. Fill a burette with this solution, and allow it to flow into a beaker containing say 15 c. c. of the volumetric solution of iodine until the brown colour of the iodine is just discharged—or, starch being added, until the blue iodide of starch is decolorized. (The latter affords the more delicate indication.) When iodine and hyposulphite of sodium react, two atoms of iodine remove two of sodium from two molecules of the hyposulphite, tetrathionate of sodium being formed, as indicated in the following equation:—



Now suppose the number of c. c. required to deoxidize the 15 c. c. of standard iodine were 14 c. c., how many c. c. of this hyposulphite solution would be equivalent to 1000 c. c. of standard iodine solution? In other words, how many c. c. would contain 24.8 grammes of hyposulphite? As 15 c. c. iodine sol. are to 14 c. c. hyposulph. sol., so are 1000 iodine sol. to x hyposulph. sol. = 933 c. c. Therefore 933 c. c. of this solution of hyposulphite would contain 24.8 grammes of the salt, and be equivalent to 1000 c. c. of the official standard solution. The 933 c. c. would be diluted to 1000 c. c. or be used without dilution. In either case its strength would, as usual, be recorded on the label. The following substances are estimated officially by means of this solution.

* "Five grains of Permanganate of Potassium dissolved in water require for decoloration a solution of forty-four grains of granulated sulphate of iron acidulated with two fluid drachms of diluted sulphuric acid."—B. P.

Solution of Chlorine.—About 10 grammes are operated upon. Excess of iodide of potassium is added—that is, to 10 grammes of solution of chlorine, about half a gramme of iodide. An amount of iodine is set free by the chlorine exactly in proportion to their atomic weights. The titration is then conducted as already described. The following show the reactions:—



It is evident, then, that 1000 c. c. of standard solution of hyposulphite of sodium, or a corresponding quantity of a solution of different strength, is equivalent to 3.55 grammes of chlorine gas. Solution of chlorine of the British Pharmacopœia contains .6 per cent. of chlorine gas.

Iodine.—Solid iodine is dissolved in solution of iodide of potassium, and titrated as already described. About .2 of a gramme is a convenient quantity to employ. 1000 c. c. of standard hyposulphite solution is equivalent, as seen in the equation, to 12.7 of iodine. The British Pharmacopœia requires "iodine" to contain 100 per cent. of real iodine. It is assumed in this operation that the iodine has been shown by qualitative analysis to be free from chlorine and bromine. These elements resemble iodine in reacting upon hyposulphite of sodium, hence would reckon as iodine in a volumetric assay.

Chlorinated Lime.—Operate on from .1 to .2 of a gramme. Dissolve in the usual quantity of water, and add excess either of dilute hydrochloric or dilute sulphuric acid and of iodide of potassium. .1 to .2 of a gramme of chlorinated lime would require .4 to .8 of a gramme of iodide of potassium. The following equations show the reactions:—



or,

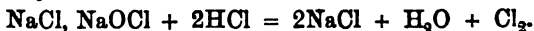


The chlorine thus set free liberates an equivalent amount of iodine, and this is titrated as before. (See the equations for

solution of chlorine.) This chlorine liberated from chlorinated lime by acids, is its available chlorine for indirect oxidizing action. It should correspond (B. P.) to 30 per cent.

Solution of Chlorinated Lime.—About 2 grammes is a convenient quantity to operate upon. 1 gramme of iodide of potassium and excess of acid should be added, and the available chlorine determined as in the case of the solid. The official (B. P.) requirement is 2.9 per cent. of available chlorine.

Solution of Chlorinated Soda.—About 2 grammes is mixed with the usual quantity of water, excess of acid added, and about 1 gramme of iodide of potassium. The available chlorine is estimated as in the case of chlorinated lime. The reaction by which the chlorine is evolved is similar:—



The action of the liberated chlorine on the iodide of potassium and the iodine on the hyposulphite solution has been given under "solution of chlorine." The official (B. P.) requirement is 2.5 per cent. of available chlorine.

QUESTIONS AND EXERCISES.

1011. For what purposes is the official volumetric solution of hyposulphite of sodium used?

1012. On what reaction is based the quantitative employment of hyposulphite of sodium?

1013. How much hyposulphite of sodium is required to show the presence of 10 parts of iodine?—*Ans.* 19.527.

1014. Calculate the amount of chlorine 4.96 parts of hyposulphite of sodium are equivalent to in volumetric analysis.—*Ans.* .71.

1015. Describe the operations included in the estimation of the strength of bleaching-powders.

1016. By what reagent is the complete absorption of free iodine by hyposulphite of sodium indicated?

MISCELLANEOUS PROBLEMS.

1017. Work sums showing how much bicarbonate of potassium is contained in an eight-ounce bottle of medicine, seven fluid drachms of which are saturated by two and a half grains of crystallized oxalic acid.—*Ans.* 36.3 grains.

1018. A sample of soda-ash is said to contain 78 per cent. of pure anhydrous carbonate of sodium: if the statement is true, how much of the official volumetric solution of oxalic acid will saturate 5 grammes of the specimen?—*Ans.* 73·6.

1019. 2·69 grammes of common brown sulphuric acid are saturated by 43·5 cubic centimètres of the official volumetric solution of soda; how much acid of 96·8 per cent. is present?—*Ans.* The 2·69 contain 2·2.

1020. Four grammes of a litre and a half of concentrated hydrocyanic acid are neutralized by 89 cubic centimètres of volumetric solution of nitrate of silver of official strength; to what volume must the bulk of the acid be diluted for the production of acid of pharmacopœial strength?—*Ans.* 9 litres.

1021. 3·18 grammes of a powder containing arsenic require for complete reaction 84 cubic centimètres of a volumetric solution of iodine, which is 1·43 per cent. weaker than the standard solution of the British Pharmacopœia; what percentage of pure arsenic is contained in the powder?—*Ans.* 12·86.

1022. How much pure metal is present in a sample of iron 1·68 gramme of which, dissolved in dilute sulphuric acid, is exactly attacked by 95·7 cubic centimètres of an official volumetric solution of red chromate of potassium which is ·6 per cent. too strong?

GRAVIMETRIC QUANTITATIVE ANALYSIS.

(For preliminary remarks on the general principles of gravimetric analysis and the relation of gravimetric and volumetric analysis to each other, see pages 608 and 609.)

ESTIMATION OF METALS.

POTASSIUM.

Outline of the Process.—This element is usually estimated in the form of double chloride of potassium and platinum. Qualitative analysis having proved the presence of potassium and other radicals in a substance, a small quantity of the material is accurately weighed, dissolved, and the other elements removed by appropriate reagents; the precipitates are well washed, in order that no trace of the potassium salt shall be lost, the resulting liquid concentrated over a water-bath (to

avoid loss that would occur mechanically during ebullition), hydrochloric acid added if necessary, solution of perchloride of platinum poured in, and evaporation continued to dryness; excess of the perchloride is then dissolved out by adding, to the dried residue, spirit of wine containing half its bulk of ether (a liquid in which the double chloride is insoluble), the mixture carefully poured on to a tared and dried filter, washed with the spirit till every trace of free perchloride of platinum is removed, the whole dried and weighed; from the resulting amount the proportion of potassium, or equivalent quantity of a salt of potassium, is ascertained by calculation.

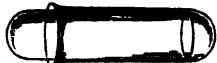
Note.—From this short description it will be seen, first, that the chemistry of quantitative is the same as that of qualitative analysis; and second, that the principle of gravimetric is the same as that of volumetric quantitative analysis:—the combining-proportions of substances being known, unknown quantities of elements may be ascertained by calculation from known quantities of their compounds.

Apparatus.—In addition to a delicate balance and weights and the common utensils, a few special instruments are used in quantitative manipulation; some of these may be prepared before proceeding with the estimation of potassium.

Filtering-paper may be of the kind known as “Swedish”—the texture of which is of the requisite degree of closeness, and its ash small in amount. A large number of circular pieces of one size, six to eight centimètres in diameter, should be cut ready for use. In delicate experiments, where a precipitate on a filter has to be ignited and the paper consequently burnt, the weight of the ash of the filter must be deducted from the weight of the residue. The ash is estimated after burning ten or twenty of the cut filters. These are folded into a small compass, a portion of a piece of platinum wire twisted a few times round the packet, so as to form a cage, the whole held by the free end of the wire over a weighed porcelain crucible placed in the centre of a sheet of glazed paper, the bundle ignited by a spirit-lamp or smokeless gas-flame, the flame allowed to impinge against the charred mass till it falls into the crucible below, any stray fragments on the

sheet carefully shaken into the crucible, the latter placed over a flame till carbon has all burnt off, and nothing but ash remains, the whole cooled, weighed, and the weight of the crucible deducted; the weight of the residue divided by the number of pieces used gives the average amount of ash in each filter.

Fig. 74.



A PAIR OF WEIGHING-TUBES.

Fig. 75.

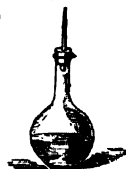


CLAMPED WATCH-GLASSES FOR WEIGHING.

A pair of *Weighing-tubes* (fig. 74), for holding dried filters during operations at the balance, may be made from two test-tubes, one fitting closely within the other. About five centimètres of the closed end of the outer and seven of the inner are cut off by leading a crack round the tube with a pencil of incandescent charcoal, and the sharp edges fused in the blow-pipe-flame. A filter, after drying, is quickly folded and placed in the narrower tube, the mouth of which is then closed by the wider tube. This prevents reabsorption of moisture from the air. A pair of *watch-glasses*, having accurately ground edges and clamped as shown in figure 75, also forms a convenient arrangement for weighing filters, etc.

The *Washing-bottle* (fig. 76), holding the *spirit of wine and ether*, is a common flask, through the cork of which a short straight tube passes. The outer end of the tube should be sufficiently narrowed to enable it to deliver a very fine stream of the liquid. The flask being inverted, the warmth of the hand expands the air and vapour to a sufficient extent to force out the liquid.

Fig. 76.



The ordinary *Washing-bottle* for quantitative operations should be formed of a flask in which water may be boiled, fitted up as usual (*vide* p. 117).

A *Water-oven* is the best form of drying-apparatus. It is a small square copper vessel, jacketed on five sides and having a door on the sixth; water is poured into the space between

the inner and outer casing, and the whole placed over a gas-lamp or other source of heat, moist air and steam escaping by appropriate apertures. Desiccation at higher temperatures than the boiling-point of water may be practised by using oil or paraffin instead of water, inserting a thermometer in the fat. The apparatus may be purchased of any maker of chemical instruments.

Pure distilled water must be used in all quantitative determinations.

Note.—In practising the operations of quantitative analysis, experiments should at first be conducted on definite salts of known composition; for the accuracy of results may then be tested by calculation.

Estimation of Potassium in the form of double chloride of potassium and platinum.—Select two or three crystals of pure nitrate of potassium, powder them in a clean mortar, dry the powder by gently heating in a porcelain crucible over a flame for a few seconds, place about a couple of décigrammes (0.2 gm.) of the powder in a counterpoised watch-glass, accurately weigh the selected quantity, transfer to a small dish, letting water from a wash-bottle flow over the watch-glass, and run into the dish, warm the dish till the nitrate is dissolved, acidulate with hydrochloric acid, add excess of aqueous solution of perchloride of platinum (a quantity containing about 0.4 of solid salt), evaporate to dryness over a water-bath. While evaporation is going on, place a filter and the weighing-tubes in the water-oven, exposing them to a temperature of 100° C. for about half an hour; fold the filter and insert it in the tubes, place them on a plate under a glass shade, and when cold accurately note their weight. Arrange the weighed filter in a funnel over a beaker. Transfer the dried and cooled platinum salt from the dish to the filter by moistening the residue with the mixture of alcohol and ether and, when the salt is loosened, pouring the contents of the dish into the paper cone. Any salt still adhering may be freed by the finger, which, together with the dish, should be washed in the stream of spirit, the rinsings at once flowing into the filter. The filtrate should have a yellowish-brown

colour, due to the excess of perchloride of platinum. If it is colourless, an insufficient amount of perchloride has been added, and the whole operation must be repeated. The washed precipitate and filter are finally dried in the water-oven, folded and placed in the weighing-tubes, the drying continued until the whole, after repeated weighing when cold, ceases to alter; the final weight is noted.

Note.—If filters are not freed from all trace of acid by thorough washing, the paper will be brittle when dry, falling to pieces on being folded.

Analytical memoranda in the note-book may have the following form :—

Watch-glass and substance	
Watch-glass	
	Substance. _____
	=====
Weighing-tubes, filter, and Pt salt . . .	
Weighing-tubes and filter	
	PtCl ₄ , 2KCl. _____
	=====

The calculations are simple :—

As $\left\{ \begin{array}{l} \text{PtCl}_4, 2\text{KCl} \\ = 489 \end{array} \right\}$ is equivalent to $\left\{ \begin{array}{l} 2\text{KNO}_3 \\ = 202 \end{array} \right\}$,
 so $\left\{ \begin{array}{l} \text{the weight of} \\ \text{double chloride} \\ \text{obtained} \end{array} \right\}$ is equivalent to x . x will be the amount

of pure nitrate of potassium in the quantity of substance operated on. x should in the present instance be identical with the weight of substance taken, because, for educational purposes, pure nitre is under examination. Only after analyses of pure substances have yielded the operator results practically identical with those by calculation, can analyses of substances of unknown degree of purity be undertaken with confidence. A table of atomic weights, from which to find molecular weights, is given in the Appendix.

Platinum residues should be preserved, and the metal recovered from them from time to time (*vide* p. 285).

Hot alcohol sometimes reduces perchloride of platinum, the metal being thrown out of solution in a finely divided form, known as *platinum black*; only aqueous solutions, therefore, of the salt should be used where heat is employed. Hence, also, in washing out excess of perchloride of platinum from the double chloride of platinum and potassium by spirit, the application of heat should be avoided.

Effervescing Potash-Water (*Liquor Potassæ Effervescens*, B. P.) is most easily estimated volumetrically (p. 651). Any adulteration by an equivalent amount of bicarbonate of sodium would, however, by that process be undetected; hence the Pharmacopœia directs that "five fluid ounces, evaporated to one fifth and 12 grains of tartaric acid added, yield a crystalline precipitate, which when dried weighs not less than 12 grains." Five fluid ounces of this preparation should contain 7.5 grains of bicarbonate, convertible into 14.1 grains of acid tartrate of potassium by 11.25 grains of tartaric acid. The method is somewhat rough, but quite efficient for "potash-water" containing nothing but bicarbonates of alkali-metals.

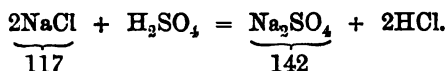
Proportional weights of equivalent quantities of potassium and its salts.

Metal	K_2	78
Oxide ("Potash")	K_2O	94
Hydrate ("Caustic Potash")	$2KHO$	112
Carbonate (anhydrous)	K_2CO_3	138
Carbonate (crystalline)	$K_2CO_3 + 16\% Aq$	164.285
Bicarbonate	$2KHCO_3$	200
Nitrate	$2KNO_3$	202
Platinum salt	$PtCl_4, 2KCl$	489

SODIUM.

Sodium is usually estimated as sulphate. Accurately weigh a porcelain crucible and lid, place within about .3 of pure rock salt, and again weigh, making a memorandum of the weights in a note-book. Add rather more strong sulphuric acid than may be considered sufficient to convert the chloride into acid sulphate of sodium. Heat the crucible gradually, the flame being first directed against the side of the crucible to avoid violent ebullition, until fumes of acid cease to be evolved, towards the end of the operation drop-

ping in one or two fragments of carbonate of ammonium to facilitate complete expulsion of all excess of acid. When cold, weigh the crucible and contents. The weight of the crucible having been deducted, the amount of sulphate obtained should be the exact equivalent of the quantity of chloride of sodium employed.



Proportional weights of equivalent quantities of sodium and its salts.

Metal	Na ₂	46
Oxide ("Soda") . . .	Na ₂ O	62
Hydrate ("Caustic Soda")	2NaHO	80
Carbonate (anhydrous) .	Na ₂ CO ₃	106
Carbonate (crystals) . .	Na ₂ CO ₃ , 10H ₂ O . .	286
Bicarbonate	2NaHCO ₃	168
Chloride	2NaCl	117
Sulphate (anhydrous) . .	Na ₂ SO ₄	142
Sulphate (crystals) . . .	Na ₂ SO ₄ , 10H ₂ O . .	322

AMMONIUM.

Salts of ammonium are, for purposes of quantitative analysis, generally converted into the double chloride of ammonium and platinum (PtCl₄·2NH₄Cl), the details of manipulation being the same as those observed in the case of potassium. About 0.15 grm. of pure, white, dry chloride of ammonium may be taken for experiment.

COMPOSITION OF THE PLATINUM SALT.

		In 1 molec. wt.	In 100 parts.
Pt	198	198	44.30
Cl ₄	35.5 × 4	213	47.64
N ₂	14.0 × 2	28	6.27
H ₈	1.0 × 8	8	1.79
		447	100.00

or, PtCl_4 . . . 340	. . . 340	. . . 76.06
$2\text{NH}_4\text{Cl}$. . . 53.5×2	. . . 107	. . . 23.94
	447	100.00

The proportion of nitrogen, ammonium, or chloride of ammonium in the double chloride may also be ascertained from the weight of platinum left on igniting the double chloride: for this purpose heat must be applied slowly, or platinum will be mechanically carried off with the gaseous products of decomposition.

Proportional weights of equivalent quantities of ammoniacal compounds.

Ammonia (gas)	2NH_3	34
Ammonium	$(\text{NH}_4)_2\text{P}$	36
Chloride of ammonium	$2\text{NH}_4\text{Cl}$	107
Platinum salt.	$\text{PtCl}_4 \cdot 2\text{NH}_4\text{Cl}$	447
"Carbonate of ammonium"	$(\text{N}_4\text{H}_{16}\text{C}_5\text{O}_8) \div 2$	118
Sulphate of ammonium	$(\text{NH}_4)_2\text{SO}_4$	132

BARIUM.

Barium is estimated in the form of anhydrous sulphate of barium (BaSO_4).

Process.—Dissolve 0.3 or 0.4 of pure crystallized and dried chloride or nitrate of barium in about half a litre of water in a beaker, heating to incipient ebullition, and slightly acidulating with hydrochloric or nitric acid. Add diluted sulphuric acid (prepared some days previously, so that sulphate of lead may have deposited) so long as a precipitate forms, keep the mixture hot for some time, set aside for half an hour, pass the supernatant liquid through a filter, gently boil the residue two or three times with more water; finally collect the precipitate on the filter, removing adherent particles from the beaker by the finger, and cleansing by a stream of hot water from the wash-bottle. The precipitate must be washed with hot water until the filtrate ceases to turn litmus-paper red, or give any cloudiness when tested with chloride of barium.

The filter and sulphate of barium, having thoroughly drained, is dried in a warm place, commonly by supporting the funnel in an inverted bottomless beaker over a sand-bath or hot plate.

The sulphate of barium is now removed from the filter, heated to drive off every trace of moisture, and weighed. This is accomplished by placing a weighed porcelain crucible (and cover) on a sheet of glazed paper, holding the filter over it, and carefully transferring the precipitate; the sides of the filter are then gently rubbed together and detached powder dropped into the crucible, the paper folded, encased in two or three coils of one end of a platinum wire and burnt over the crucible, ash and any particles on the sheet of paper dropped into the sulphate of barium, the open crucible exposed over a flame till its contents are quite white, covered, cooled, and weighed.

	Formulae.	Molecular weights.
Chloride of barium . . .	$\text{BaCl}_2, 2\text{H}_2\text{O}$. . .	244
Nitrate of barium . . .	$\text{Ba}2\text{NO}_3$. . .	261
Sulphate of barium . . .	BaSO_4 . . .	233

Composition of sulphate of barium.

		In 1 molec. wt.	In 100 parts.
Ba	137	137 . . .	58·80
S	32	32 . . .	13·73
O ₄	16×4	64 . . .	27·47
		233	100·00

In these educational experiments it is unnecessary to take filter-ash into account. Inevitable mistakes of manipulation commonly cause far greater errors.

CALCIUM.

Calcium is usually thrown out of solution in the form of oxalate, the precipitate ignited, and the resulting carbonate weighed.

Process.—Dissolve 0·3 or 0·4 of dried colourless crystals of calc-spar in about a third of a litre of water acidulated with

hydrochloric acid, heat the solution to near the boiling-point, add excess of solution of oxalate of ammonium, then ammonia until, after stirring, the liquid smells strongly ammoniacal: set aside in a warm place for twelve hours. Carefully pour off the supernatant liquid, passing it through a filter; add hot water to the precipitate, set aside for half an hour, again decant, and, after once more washing, transfer the precipitate to the filter, allowing all contained fluid to pass through before a fresh portion is added. Wash the precipitate with hot water, avoiding a rapid stream, or the precipitate may be driven through the pores of the paper. Dry, transfer to a weighed crucible, and incinerate, as described for sulphate of barium, and slowly heat the precipitate till the bottom of the crucible is just visibly red when seen in the dark. As soon as the residue is white, or only faintly grey, remove the lamp, cool, and weigh.

The resulting carbonate of calcium should have the same weight as the calc-spar from which it was obtained. If loss has occurred, carbonic acid gas has probably escaped. In that case moisten the residue with water, and after a few minutes test the liquid with red litmus or turmeric paper; if an alkaline reaction is noticed, it is due to the presence of caustic lime. Add a small lump of carbonate of ammonium, evaporate to dryness over a water-bath, and again ignite, this time being careful not to go beyond the prescribed temperature. The treatment may, if necessary, be repeated.

Proportional weights of equivalent quantities of calcium and its salts.

Metal	Ca	40
Oxide (quicklime). . .	CaO	56
Hydrate (slaked lime)	Ca ₂ HO	74
Carbonate	CaCO ₃	100
Sulphate (anhydrous)	CaSO ₄	136
Sulphate (crystalline or precipitated) . . .	CaSO ₄ , 2H ₂ O	172
Chloride	CaCl ₂	111
Phosphate (of bones).	(Ca ₃ 2PO ₄)310 ÷ 8 . .	103·3

MAGNESIUM.

Process 1.—The light or heavy carbonate of magnesium of pharmacy may be estimated by heating a weighed quantity to redness in a porcelain crucible. If it has the composition indicated by the formula given in the British Pharmacopœia ($3\text{MgCO}_3, \text{Mg}_2\text{HO}, 4\text{H}_2\text{O}$), it will yield 42 per cent. of magnesia (MgO). According to that work, the purity of even sulphate of magnesium ($\text{MgSO}_4, 7\text{H}_2\text{O}$) may be determined by boiling a weighed quantity with excess of carbonate of sodium, collecting the precipitate, washing, drying, igniting, and weighing the resulting magnesia (MgO). The crystallized sulphate should afford 16.26 per cent. of oxide. The official solution of carbonate of magnesium in carbonic acid water (*Liquor Magnesice Carbonatis*, B. P.) should yield five grains of pure oxide of magnesium per fluid ounce.

Process 2.—The general form in which magnesium is precipitated is as phosphate of ammonium and magnesium ($\text{MgNH}_4\text{PO}_4, 6\text{H}_2\text{O}$); this, by heat, is converted into pyrophosphate of magnesium ($\text{Mg}_2\text{P}_2\text{O}_7$). Accurately weigh a small quantity (0.4 to 0.5) of pure dry crystals of sulphate of magnesium, dissolve in two to three hundred cubic centimetres of cold water in a beaker, add chloride of ammonium, ammonia, and phosphate of sodium or ammonium, agitate with a glass rod (without touching the sides of the vessel, or crystals will firmly adhere to the rubbed portions), and set aside for twelve hours. Collect on a filter, wash the precipitate with water containing a tenth of its volume of the strongest solution of ammonia, until the filtrate ceases to give a precipitate with an acidulated solution of nitrate of silver. Dry, transfer to a crucible, burn the filter in the usual way, heat slowly to redness, cool, and weigh.

Proportional weights of equivalent quantities of magnesium salts.

Pyrophosphate.	$\text{Mg}_2\text{P}_2\text{O}_7$	222
Sulphate.	$2(\text{MgSO}_4, 7\text{H}_2\text{O})$	492
Oxide.	$2(\text{MgO})$	80
Official carbonate.	$(3\text{MgCO}_3, \text{Mg}_2\text{HO}, 4\text{H}_2\text{O}) \div 2$	191

ZINC.

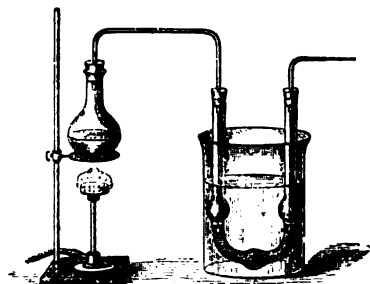
Zinc is usually estimated as oxide (ZnO), occasionally as sulphide (ZnS).

Process.—Dissolve a weighed quantity (0.5 to 0.6) of sulphate of zinc in about half a litre of water in a beaker, heat to near the boiling-point, add carbonate of sodium in slight excess, boil, set aside for a short time; pass the supernatant liquid through a filter, gently boil the precipitate with more water, again decant; repeat these operations two or three times; collect the precipitate on the filter, wash, dry, transfer to a crucible, incinerate, ignite, cool, and weigh. 287 (= molec. weight) of sulphate should yield 81 (= molec. weight) of oxide.

MANGANESE.

To ascertain its value for evolving chlorine from hydrochloric acid, a weighed quantity of finely powdered black

Fig. 77.



oxide of manganese is heated in a small flask with pure hydrochloric acid, and the resulting chlorine conveyed into a U-tube containing solution of iodide of potassium. The amount of iodine thus freed is estimated by the volumetric solution of hyposulphite of sodium. 127 of iodine indicate 35.5 of chlorine.

Manganese may also be estimated by the reaction and apparatus described under "Oxalates," p. 704.

ALUMINIUM.

Aluminium is always precipitated as hydrate ($\text{Al}_2\text{6HO}$) and weighed as oxide (Al_2O_3).

Process.—Dissolve about two grammes of pure dry ammonium-alum in half a litre of water, heat the solution, add chloride of ammonium and a slight excess of ammonia, boil gently till the odour of ammonia has nearly disappeared, set aside for the hydrate to deposit, pass the supernatant liquid through a filter, wash the precipitate three or four times by decantation, transfer to the filter, finish the washing, dry, burn the filter, ignite in a covered crucible, and weigh.

$\text{Al}_2\text{3SO}_4, (\text{NH}_4)_2\text{SO}_4, 24\text{H}_2\text{O}$ 907

Al_2O_3 103

Per cent. of Al_2O_3 yielded by ammonium-alum 11·356

QUESTIONS AND EXERCISES.

1023. Give details of the manipulations observed in gravimetrically estimating salts of potassium or of ammonium.

1024. What quantity of chloride of sodium is contained in a sample of rock-salt 0·351 gramme of which yields 0·426 of sulphate of sodium?—*Ans.* 100 per cent. (It is absolutely pure.)

1025. To what amount of the official alum is 0·894 of a gramme of the double chloride of platinum and ammonium equivalent?—*Ans.* 1·814 gramme.

1026. Find the weight of sulphate of barium obtainable from 0·522 of nitrate.—*Ans.* 0·466.

1027. Describe the usual method by which salts of calcium are estimated.

1028. By what quantitative processes may the official salts of magnesium be analysed?

1029. Calculate the proportion of pure sulphate of zinc in a sample of crystals 0·574 of which yield 0·161 of oxide.—*Ans.* 99·3 per cent.

1030. Ascertain the weight of alumina (Al_2O_3) which should be obtained from 1·814 gramme of ammonium-alum.

IRON.

Iron and its salts are gravimetrically estimated in the form of ferric oxide (Fe_2O_3).

Compounds containing organic acidulous radicals are simply incinerated, and the resulting oxide weighed. Thus 1 gramme of the official citrate of iron and ammonium (*Ferri et Ammoniae Citras*, B. P.) incinerated, with exposure to air, leaves not less than .27 of ferric oxide. A small quantity of the salt is weighed in a tared covered porcelain crucible, flame cautiously applied until vapours cease to be evolved, the lid then removed, the crucible slightly inclined and exposed to a red heat until all carbonaceous matter has disappeared. The residual ferric oxide is then weighed. The tartrate of potassium and iron (*Ferrum Tartaratum*, B. P.) is treated in the same manner, except that the ash must be washed and again heated before weighing, in order to remove carbonate of potassium produced during incineration; 5 grammes should yield 1.5 gramme of ferric oxide.

From other compounds of iron, soluble in water or acid, the metal is precipitated in the form of hydrate (Fe_26HO) by solution of ammonia, and converted into oxide (Fe_2O_3) by ignition. Dissolve a piece (about 0.2) of the purest iron obtainable (piano-wire), accurately weighed, in water acidulated with hydrochloric acid; add a few drops of nitric acid and gently boil; pour in excess of ammonia, stir, set aside till the ferric hydrate has deposited, pass the supernatant liquid through a filter, treat the precipitate three or four times with boiling water; transfer to the filter, wash till the filtrate yields no trace of chlorine (for chloride of ammonium will decompose ignited ferric oxide, with volatilization of ferric chloride), dry, and ignite as usual, and weigh. Iron in the official solutions (*Liquor Ferri Perchloridi Fortior*, *Liquor Ferri Pernitratis*, and *Liquor Ferri Persulphatis*) is estimated by this general process.

The proportion of metallic iron in a mixture of iron and oxides of iron may be determined by digestion in a strong

solution of iodine in iodide of potassium, which attacks the metal only. The reduced iron of pharmacy (*Ferrum Redactum*, B. P.) is in good condition so long as it contains, as shown by this method, half its weight of free metal.

Proportional weights of equivalent quantities of iron and its salts.

Metal	Fe_2	112
Ferric oxide . . .	Fe_2O_3	160
Ferric hydrate . .	Fe_36HO	214
Ferric chloride . .	Fe_2Cl_6	325
Ferric sulphate . .	Fe_23SO_4	400
Ferrous sulphate .	$2(\text{FeSO}_4, 7\text{H}_2\text{O})$. .	556

ARSENICUM.

Arsenic (As_2O_3) is usually estimated volumetrically (*vide* p. 664). With certain precautions arsenicum may also be precipitated and weighed as sulphide (As_2S_3).

Process 1.—The pure, white, massive arsenic (about 0·2) is dissolved in a flask in a small quantity of water containing bicarbonate of sodium or potassium, the liquid being heated. A slight excess of hydrochloric acid is then added, and sulphuretted hydrogen gas passed through the solution so long as a precipitate falls, the mouth of the flask being stopped by a plug of cotton-wool (to prevent undue access of air and consequent decomposition of the gas, resulting in precipitation of sulphur). The mixture is warmed in the flask and carbonic acid gas passed through it until the odour of sulphuretted hydrogen has nearly disappeared; the precipitate is collected on a tared filter, washed as quickly as possible with hot water containing a little sulphuretted hydrogen, dried in a water-oven and weighed. 198 parts of arsenic should yield 246 of sulphide of arsenicum.

Process 2.—The arsenicum must be present in the arsenic condition. If the operator is not certain that this is the case, the solution must be warmed with a little hydrochloric acid and a few grains of chlorate of potassium added until a distinct odour of chlorous vapour is evolved—which is then allowed

to escape by continued application of heat. To the solution thus obtained ammonia, which must produce no turbidity, is added in excess, and then magnesia mixture (*see* p. 705). The solution is set aside for 24 or 48 hours. The precipitate is collected on a filter and washed with as little ammonia water (1 to 3) as possible until the filtrate ceases to give a reaction for chlorides. The precipitate is then dried on the filter, the precipitate and filter paper burned, and the whole gently ignited in a crucible, and weighed. The residue is represented by the formula $Mg_3As_2O_7$.

ANTIMONY.

The metal is precipitated in the form of sulphide (Sb_2S_3), with the precautions observed in estimating arsenicum—a small quantity of tartaric acid, as well as hydrochloric, being added, to prevent the precipitation of an oxysalt. If the sulphuretted hydrogen be passed through a hot solution, the particles of precipitate aggregate better and the latter may be more quickly filtered out and washed. The experiment may be performed on about half a gramme of pure tartar-emetic: the salt should yield nearly half its weight (49.56 per cent.) of sulphide. According to Fresenius, the sulphide dried at $100^\circ C$. still contains 2 per cent. of water, and must be heated in a current of carbonic acid gas, until it turns from an orange to a black colour, before all moisture is expelled. In the British Pharmacopœia the purity of tartar-emetic (*Antimonium Tartaratum*), and the strength of solution of chloride of antimony (*Liquor Antimonii Chloridi*), are determined by the above process.

COPPER.

Copper is precipitated from its solutions and weighed either (1) as metal (Cu_2), or (2) as oxide (CuO).

Process 1.—Dissolve about half a gramme of dry crystallized sulphate of copper in a small quantity of water, in a tared porcelain crucible or beaker, acidulate with hydrochloric acid, introduce a fragment or two of pure zinc, cover

the vessel with a watch-glass, and set aside till evolution of hydrogen has ceased and the still acid liquid is colourless. The copper is then washed with hot water by decantation until no trace of acid remains, the precipitate drained, rinsed with strong spirit of wine, dried in the water-oven, and weighed.

Process 2.—From a solution acidulated by sulphuric acid and placed in a platinum crucible, copper may be entirely deposited in a coherent form by a weak current of electricity, the crucible being connected with the zinc pole of the battery, a platinum spatula suspended in the solution forming the positive pole. The crucible may afterwards be freed from the deposited copper by nitric acid.

Process 3.—About three-fourths of a gramme of sulphate of copper is accurately weighed, dissolved in half a litre of water, the liquid boiled; dilute solution of potash or soda is then added till no more precipitate falls, ebullition continued for a short time, and the beaker set aside; the supernatant liquid is decanted, the precipitate boiled with water twice or thrice, collected on a filter, washed, dried, transferred to a crucible, the filter incinerated, and its ash moistened with a drop of nitric acid; the whole is finally heated strongly, cooled, and weighed.

249·5 parts of sulphate of copper yield 79·5 of oxide, or 63·5 of metal.

Other processes.—*Vide Pharmaceutical Journal* for April 3, 1880, p. 801.

BISMUTH.

Dissolve 0·3 or 0·4 of pure oxycarbonate of bismuth ($2\text{Bi}_2\text{O}_3\text{CO}_3, \text{H}_2\text{O}$) (*Bismuthi Carbonas*, B. P.) in a small quantity of hydrochloric acid, dilute with water slightly acidulated by hydrochloric acid, pass excess of sulphuretted hydrogen through the liquid, collect the precipitate on a tared filter, wash, dry at 100°C ., and weigh. The sulphide must not be exposed too long in the water-oven, or it will increase in weight owing to absorption of oxygen; hence it should be tested in the balance every half-hour during desic-

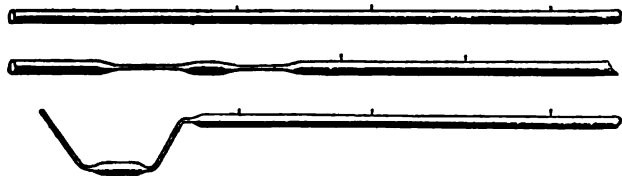
cation. 517 of oxycarbonate should yield 512 of sulphide (Bi_2S_3). The strength of the official solution of citrate of bismuth and ammonium (*Liquor Bismuthi et Ammonie Citratis*, B. P.) is determined by this process. "Three fluid drachms of the solution, mixed with an ounce of distilled water, and treated with sulphuretted hydrogen in excess, yield a black precipitate, which, collected, washed, and dried, weighs 9.92 grains. One fluid drachm yields three grains of oxide of bismuth." The atomic weight of bismuth is 208.

MERCURY.

This element may be (1) isolated and estimated in the form of metal, or precipitated and weighed as (2) mercurous chloride, or (3) mercuric sulphide.

Process 1.—The process by which the metal itself is separated is one of distillation into a bulb surrounded by water. About half a mètre of the difficultly fusible German glass known as *combustion-tubing* is sealed at one end after the manner of a test-tube (fig. 78); a mixture of bicarbonate of

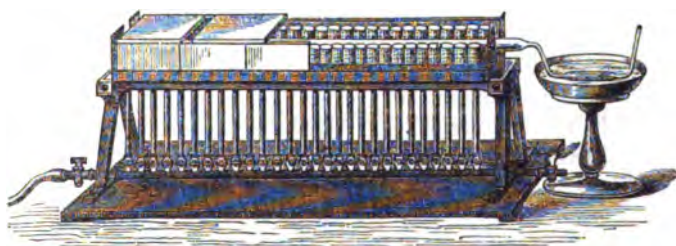
Figs. 78, 79, 80.



sodium and dry chalk is then dropped into the tube to the height of two or three centimètres, and, next, several small fragments of quicklime so as to occupy another centimètre: a mixture of about a gramme of pure calomel or corrosive sublimate with enough powdered quicklime to occupy 10 or 12 centimètres of the tube is added, then the lime-rinsings of the mixing-mortar, a layer of a few centimètres of powdered quicklime, and finally a plug of *asbestos*. The whole should occupy two-thirds of the tube. The part of the tube just

above the asbestos is now softened in the blowpipe-flame and drawn out about a decimètre to the diameter of a narrow quill (fig. 79); again drawn out to the same extent at a point two or three centimètres nearer the mouth (fig. 79), and any excess of tubing cut off. The bulb thus formed may be enlarged by softening and blowing. The tube is next softened at a point close to but anterior to the asbestos, and bent to form an obtuse angle; the tube is then softened close to the bulb and slightly bent so that the bulb may be parallel with the large tube; then softened on the other side of the bulb, and the terminal tube bent to an obtuse angle, so that, the tube being held in a horizontal position, the bulb may be sunk in water, and the terminal tube point upwards (fig. 80). The long tube is now laid in the gas-furnace found in most laboratories (fig. 81), a basin so placed that the bulb of the apparatus may be cooled by being surrounded by water, the part of the tube occupied by asbestos heated to redness, and the flame slowly lengthened until the whole tube is red-hot.

Fig. 81.



DISTILLATION OF MERCURY FOR QUANTITATIVE PURPOSES.

Under the circumstances just described the mercurial compound volatilizes, is decomposed by the lime, and its acidulous radical fixed, the mercury carried in vapour to and condensed in the bulb, the carbonic acid gas evolved from the bicarbonate of sodium and chalk washing out the last portions of mercury-vapour from the tube. When the distillation is considered to be complete, the dish of water is removed, the

bulb dried, and then detached by help of a file at a point beyond any sublimate of mercury. The bulb is lastly weighed, the mercury shaken, or dissolved out, and the tube again dried and weighed. The difference between the weights gives the weight of the mercury.

Process 2.—The process by which mercury is separated in the form of calomel, consists in adding solution of hydrochloric and of phosphorous acids to an aqueous or even acid solution of a weighed quantity of the mercurial compound, setting the mixture aside for twelve hours, collecting the precipitate on a tared filter, washing, drying at 100°C ., and weighing (Rose). The experiment may be tried on half a gramme to a gramme of corrosive sublimate.

Process 3.—Two or three décigrammes of corrosive sublimate are dissolved in water, the solution acidulated with hydrochloric acid, excess of sulphuretted hydrogen passed through it, the precipitate collected on a tared filter, washed with cold water, dried at 100°C ., and weighed.

Proportional weights of equivalent quantities of mercury and its salts.

Metal	Hg.	200
Mercurous chloride . .	HgCl	235.5
Mercuric chloride . .	HgCl ₂	271
Mercuric sulphide . .	HgS	232

LEAD.

Lead is generally estimated either as (1) oxide, (2) sulphate, (3) chromate, or (4) metal.

Process 1.—Weigh out one or two grammes of pure acetate of lead in a covered crucible, previously tared, and heat slowly until no more vapours are evolved. Remove the lid, stir down the carbonaceous mass with a clean iron wire, and keep the crucible in the flame so long as any carbon remains unconsumed. Introduce some fragments of fused nitrate of ammonium, and again ignite until no metallic lead remains.

and all excess of the nitrate has been decomposed. Cool and weigh the resulting oxide (PbO).

Process 2.—Dissolve 0.4 or 0.5 of a gramme of acetate of lead in a small quantity of water, drop in diluted sulphuric acid, add to the mixture twice its bulk of methylated spirit of wine, and set aside. Decant the supernatant liquid, collect the sulphate on a filter, wash with spirit, dry, transfer to a porcelain crucible, removing as much of the sulphate as possible from the paper, incinerate on the crucible-lid (not in a platinum coil, for the particles of reduced lead would unite with the platinum by fusion), ignite, cool, and weigh.

Process 3.—About half a gramme of acetate of lead is dissolved in two or three hundred c. c. of water, acetic acid added, and then solution of red chromate of potassium. Collect the precipitate on a tared filter, wash, dry at 100°C ., and weigh.

Process 4.—In certain cases, notably in that of commercial "white lead," the lead may be estimated in the metallic state by means of cyanide of potassium. The lead paint (about 20 grammes) is weighed and carefully incinerated. The residue, a mixture of metallic lead and oxide of lead, is then mixed with several times its bulk of cyanide of potassium and the whole heated to fusion. With careful manipulation the lead collects in one globule, which, after cooling, may readily be separated from the mixed cyanide and cyanate and weighed. Commercially pure white lead should yield 74 per cent. of lead.

Molecular weights of salts of lead.

Metal . . .	Pb.	207
Acetate . . .	$\text{Pb2C}_2\text{H}_3\text{O}_2, 3\text{H}_2\text{O}$. .	379
Oxide . . .	PbO	223
Sulphate . .	PbSO_4	303
Chromate . .	PbCrO_4	323.5

SILVER.

Compounds of silver which are readily decomposed by heat are estimated in the form of (1) metal, others usually as (2) chloride (AgCl), but sometimes as (3) cyanide (AgCN).

Process 1.—Heat about a gramme of oxide of silver (Ag_2O) in a tared crucible, cool and weigh. 232 of oxide yield 216 of metal. “29 grains heated to redness yield 27 grains of metallic silver.”—*Brit. Pharm.*

Process 2.—Dissolve 0.4 or 0.5 of pure dry crystals of nitrate of silver in water, acidulate with two or three drops of nitric acid, slowly add hydrochloric acid, stirring rapidly, until no more precipitate falls. Pour off the supernatant liquid through a filter, wash the chloride of silver once or twice with hot water, transfer to the filter, complete the washing, and dry. After removing as much as possible of the precipitate from the paper to the crucible, burn the filter, letting its ash fall on the inverted lid of the crucible, moisten with a drop of nitric acid, warm, add a drop of hydrochloric acid, evaporate to dryness, replace the lid on the crucible, ignite the whole until the edges of the mass of chloride begin to fuse; cool and weigh. 170 of nitrate yield 143.5 of chloride. According to the British Pharmacopœia, 10 parts of nitrate should thus yield 8.44 of chloride, and the filtrate from the chloride evaporated to dryness should leave no residue, indicating absence of nitrates of potassium or sodium and other similar foreign substances.

Process 3.—Cyanide of silver may be collected on a tared filter and dried at 100°C 170 of nitrate yield 134 of cyanide.

Silver and its salts may be volumetrically estimated by a standard solution of chloride of sodium.

Cupellation.—The amount of silver in an alloy may be also determined by a dry method. The metal is folded in a piece of thin sheet lead, placed on a *cupel* (*cupella*, little cup, made of compressed bone-earth) and heated in a furnace, the cupel being protected from the direct action of the flame by a muff-shaped or, rather, oven-shaped case termed a muffle. The metals melt, the baser become oxidized, the oxide of lead fusing and dissolving the other oxides; the fluid oxides are absorbed by the porous cupel, a button of pure silver remaining. An alloy suspected to contain 95 per cent. of silver requires about three times its weight of lead for successful

cupellation; if $92\frac{1}{2}$ per cent. (English silver coin), between five and six times its weight of lead is necessary.

QUESTIONS AND EXERCISES.

1031. Explain the gravimetric process by which the strength of the official solutions of ferric chloride, nitrate, and sulphate are determined.

1032. Mention the various amounts of ferrous and ferric salts equivalent to 100 parts of metal.

1033. State the precautions necessary to be observed in estimating arsenicum or antimony in the form of sulphide.

1034. In what form are the official compounds of bismuth weighed for quantitative purposes?

1035. Give an outline of the process by which mercury may be isolated from its official preparations and weighed in the metallic condition.

1036. Describe three methods for the quantitative analysis of salts of lead; and the weights of the respective precipitates, supposing 0.56 of crystallized acetate to have been operated on in each case.

1037. Describe the processes by which silver is estimated in the forms of metal, chloride, and cyanide.

1038. What proportions of nitrate of silver are indicated, respectively, by 15 of metal, 9.8 of chloride, and 8.1 of cyanide?

1039. Describe cupellation.

GRAVIMETRIC ESTIMATION OF THE ACIDULOUS RADICALS OF SALTS.

CHLORIDES.

Free chlorine (chlorine-water) and compounds which by action of acids yield free chlorine (Chlorinated Lime, Chlorinated Soda, and their official Solutions) are estimated volumetrically by a standard solution of hyposulphite of sodium (*vide* p. 671). The amount of combined chlorine in pure chlorides (HCl, NaCl) may also be determined by volumetric analysis with a standard solution of nitrate of silver (p. 659).

Combined chlorine is gravimetrically estimated in the form of chloride of silver, the operation being identical with that just described for silver salts (p. 694). 58.5 parts of pure, colourless, crystallized chloride of sodium (rock-salt) yield 143.5 of chloride of silver.

IODIDES.

Free iodine is estimated volumetrically by solution of hyposulphite of sodium (*vide* p. 671).

Combined iodine is determined gravimetrically in the form of iodide of silver, the operations being conducted as with chloride of silver. Iodide of potassium may be used for an experimental determination: $KI = 166$ should yield $AgI = 235$. Of the official iodide of cadmium (*Cadmii Iodidum*, B. P.) it is stated that "ten grains dissolved in water, and nitrate of silver added in excess, give a precipitate which, when washed with water and afterwards with half an ounce of solution of ammonia, and dried, weighs 12.5 grains."

In presence of chlorides and bromides the iodine in iodides may be precipitated and weighed as iodide of palladium.

Moisture in iodine is estimated by loss on exposing a weighed quantity of iodine in a capsule over a dish of sulphuric acid under a small bell jar: or by adding to a weighed sample five or six times as much mercury, or twice as much zinc, and a little water, drying and weighing. The product is the amount of metal employed plus that of the *dry* iodine in the sample.

BROMIDES.

Free bromine may be estimated by shaking with excess of solution of iodide of potassium, and then determining the equivalent quantity of liberated iodine by a standard solution of hyposulphite of sodium (p. 671).

The bromine in bromides may be precipitated and weighed as bromide of silver, the manipulations being the same as those for chloride of silver: 0.2 to 0.3 of pure bromide of potassium may be used for an experimental analysis.

CYANIDES.

The *hydrogen cyanide* (hydrocyanic acid) is usually estimated volumetrically (*vide* p. 659).

From all soluble cyanides cyanogen may be precipitated by nitrate of silver, after acidulating with nitric acid, the cyanide of silver collected on a tared filter, dried at 100° C. and weighed.

Of the official Diluted Hydrocyanic Acid, it is stated that one hundred grains (or 110 minims) precipitated by solution of nitrate of silver yield ten grains of dry cyanide of silver.

Cyanide of silver.

		In 1 molec. wt.	In 100 parts.
Silver . .	Ag . .	107.93 . .	80.59
Cyanogen .	CN . .	26.00 . .	19.41
		133.93	100.00

NITRATES.

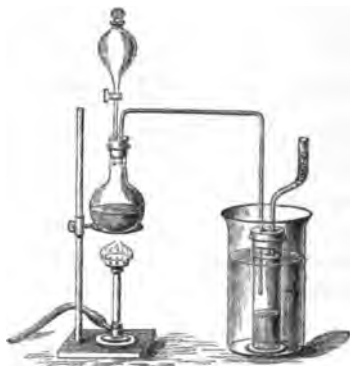
Nitrates cannot be estimated by direct gravimetric analysis, none of the basylous radicals yielding a definite nitrate insoluble in water. With some difficulty they may be determined by indirect volumetric methods.

Process.—The following (Thorpe's) method depends upon the fact (Gladstone and Tribe) that when zinc upon which copper is deposited in a spongy form is boiled with water, hydrogen is evolved. Thorpe found that in a solution containing nitrates the nascent hydrogen converts the whole of the nitrogen of the nitrates into ammonia, which may be collected and estimated. (The oxygen of the nitrate is simultaneously converted into water, the nitrate-metal into hydrate, and the zinc into hydrate of zinc. The power of the copper-zinc couple is considered to depend largely on the hydrogen absorbed by the finely divided metal.)

An apparatus such as shown in fig. 82 should be constructed. A flask (about 100 c. c.) is fitted with a clean sound cork perforated for a delivery-tube, which should be of strong

glass tubing of about quarter-inch bore, and for a stoppered funnel, which should have about half the capacity of the flask. The whole is supported by a clamp or on wire-gauze.

Fig. 82.



ESTIMATION OF NITRATES.

The outer jar shown in the figure should have a capacity of two or three litres, and the inner receiving-jar should be capable of holding 200 c. c. The latter is fitted with a cork perforated for the delivery-tube, and perhaps for another tube containing fragments of glass moistened with acidulated water to prevent possible loss of ammonia—though the latter tube is practically found to be almost unnecessary. The addition of washing-bottle tubes is also recommended (Stapleton) as convenient for obtaining the distillate from the jar without dismounting the apparatus.

A few strips of clean zinc (*granulated* zinc recently cleansed with dilute acid—Dunstan), are boiled in a beaker with a four per cent. solution of sulphate of copper, the operation being repeated with a fresh portion of solution until an adherent and fairly thick coating of finely divided copper is deposited. The pieces of metal are well washed and introduced into the flask, which is then half filled with pure water. To avoid transference, the flask itself may be used

instead of the beaker. The funnel also of the apparatus is filled with pure water. Water is now placed around the inner receiver in the outer jar, and, the connections being sound, heat is applied with the view of freeing the apparatus itself from any trace of ammonia. When the contents of the flask are evaporated nearly to dryness, pure water is admitted from the funnel until the flask is again about half full (the funnel should be filled again at once), and the distillation carried on as before. This must be repeated until no further trace of ammonia is evolved, when the apparatus is ready for use. On each occasion that the apparatus is used it must be freed from ammonia in this way. A suitable quantity of the substance to be estimated is now introduced (in the case of potable waters the prepared solid residue from 100 c. c.) and water added, if necessary, until the flask is half full. Heat is now applied and the operation conducted in the manner already described until ammonia ceases to come over—a point which usually occurs in the case of water-residues when the flask has been twice or thrice charged with water and the distillate is about 100 c.c. The warm water from the upper part of the cooling-jar may be removed by a siphon or otherwise, cold water being introduced from time to time.

The ammonia being all evolved, disconnect the flask and receiver simultaneously (unless washing-bottle tubes are fitted), and treat the contents of the latter by the Nessler method described on page 648.—Urea yields but traces of ammonia by this process; and neither the sulphates nor chlorides of the alkali-metals affect the result.—The method is only applicable to highly dilute solutions of nitrates, for with stronger solutions oxides of nitrogen are formed and escape.

Another process (Pelouze's, improved by Fresenius) consists in adding the nitrate to an acid solution of a ferrous salt of known strength, and, when reaction is complete, estimating the amount of ferrous salt unattacked by volumetric solution of red chromate, or of permanganate. Three molecular weights of converted ferrous salt indicate one molecular weight of nitric acid. Regeneration of nitric or nitrous acids

by aërial oxidation of the nitric oxide evolved is prevented either by a current of carbonic acid gas, or by using a closed flask in which is a *Bunsen-valve* (i.e. a short attached piece of india-rubber tubing closed at the free extremity, and having a sharp longitudinal slit in it a third of an inch long—a slit by which gases can escape but cannot re-enter).

SULPHIDES.

Process 1.—Soluble sulphides (H_2S , NaHS , e.g.) may be estimated volumetrically by adding to the aqueous liquid a measured excess of an alkaline solution of arsenic of known strength, neutralizing by hydrochloric acid, diluting to any given volume, filtering off the sulphide of arsenicum precipitated, taking a portion of the filtrate equal to a half or a third of the original volume, and, after neutralizing by acid carbonate of sodium, estimating the residual arsenic by the standard iodine solution (*vide* p. 664). The process may be tried on a measured volume of sulphuretted hydrogen (the weight of which is easily calculated; 1 litre of hydrogen = 0.0896 gramme) absorbed by a strong solution of soda or potash.

Process 2.—Sulphur and sulphides may also be quantitatively analysed by oxidizing to sulphuric acid and precipitating in the form of sulphate of barium. A couple of décigrammes of a pure metallic sulphide may be decomposed by careful deflagration with a mixture of chlorate of potassium and carbonate of sodium, the product dissolved in water, acidulated with hydrochloric acid, solution of chloride of barium added, and the precipitated sulphate of barium purified and collected as described in connection with the estimation of barium (p. 680). Many sulphides may be oxidized in a flask by chlorate of potassium and hydrochloric acid, and then precipitated by chloride of barium. Experimental determinations may also be made on a weighed fragment of sulphur, about 0.1, cautiously fused with a little solid caustic alkali, and the product oxidized while hot by the slow addition of powdered nitrate or chlorate of potassium, or, when cold, by treatment

with chlorate of potassium and hydrochloric acid, and subsequent precipitation by chloride of barium.

Note.—Fusions performed by help of a gas-lamp must be carefully conducted; for any alkali that may creep over the side of a crucible will certainly absorb sulphurous acid from the products of combustion of the gas, and error result.

Process 3.—Soluble sulphides may also be treated with excess of an alkaline arsenite, arsenious sulphide then precipitated by the addition of hydrochloric acid, and the precipitate collected and weighed with the usual precautions (*vide* p. 687).

Weights of equivalent quantities of sulphur and its compounds.

Sulphur	S	32
Sulphuretted hydrogen . .	H ₂ S	34
Sulphate of barium . .	BaSO ₄	233
Arsenious sulphide . . .	(As ₂ S ₃) ÷ 3	82
Bisulphide of iron . . .	(FeS ₂) ÷ 2	60
Sulphide of lead	PbS	239

SULPHITES.

Sulphites are usually estimated volumetrically by a standard solution of iodine (*vide* p. 663). Sulphites insoluble in water are diffused in that menstruum, hydrochloric acid added, and the iodine solution then dropped in.

If necessary, sulphites may be estimated gravimetrically by oxidation and precipitation in the form of sulphate of barium.

SULPHATES.

These salts are always precipitated and weighed as sulphate of barium, the manipulations being identical with those performed in the determination of barium by means of sulphates (*vide* p. 680). The purity of Sulphate of Sodium (*Sodæ Sulphas*, B. P.), and the presence of not more than a given amount of sulphuric acid in Vinegar (*Acetum*, B. P.), are directed, in the British Pharmacopœia, to be ascertained by this process. Ten grains of sulphate of sodium yield 7·236 of

sulphate of barium. Five ounces of vinegar should yield not more than about one-third of a gramme of sulphate of barium.

The amount of free sulphuric acid or hydrochloric acid in vinegar, lemon juice, lime juice, etc., may also be ascertained volumetrically by adding a known quantity of standard solution of soda, evaporating to dryness, incinerating, dissolving in water, and by standard acid estimating the quantity of soda still remaining free. The soda lost indicates the amount of free mineral acid (Hegner). Thresh first estimates the chlorine in a sample of vinegar, then adds a known additional amount of chlorine, preferably in the form of chloride of barium, evaporates, ignites, treats with water, adds bicarbonate of sodium to remove excess of barium, filters, and again estimates the chlorine. A loss of 71 of chlorine (Cl_2) indicates 98 of free sulphuric acid (H_2SO_4).

The method of estimating free sulphuric, nitric, or hydrochloric acids, proposed by Spence and Esilman, is founded on their power of decolourizing a standard solution of ferric acetate.

Proportional weights of equivalent quantities of sulphates.

The sulphuric radical . . .	SO_4	96
Sulphuric acid	H_2SO_4	98
Sulphate of barium	BaSO_4	233

CARBONATES.

Carbonates are usually estimated by the loss in weight they undergo on the addition of a strong acid.

Process 1.—A small light flask is selected—of such a size that it can be conveniently weighed in a delicate balance. Two narrow glass tubes are fitted to the flask by a cork—the one straight, extending from about two or three centimètres above the cork to the bottom of the flask,



the other cut off close to the cork on the inside and curved outwards so as to carry a thin drying-tube horizontally above the flask (*see* fig. 83). The drying-tube is nearly filled with small pieces of chloride of calcium, a plug of cotton-wool preventing escape of any fragments at either end, and is attached by a pierced cork to the free extremity of the curved tube of the flask. A weighed quantity of any pure soluble carbonate is placed in the flask, a little water added, a miniature test-tube containing sulphuric acid lowered into the flask by a thread and supported so that the acid may not flow out, the cork inserted, the outer end of the piece of the straight glass tubing closed by a fragment of cork or wax, and the whole weighed. The apparatus is then inclined so that the oil of vitriol and carbonate may slowly react; carbonic acid gas is evolved and escapes through the horizontal tube, any moisture being retained by the chloride of calcium. When effervescence has ceased, the gas still remaining in the vessel is sucked out; this is accomplished by adapting a piece of india-rubber tubing to the end of the drying-tube, removing the small plug from the straight tube, and aspirating slowly with the mouth for a few minutes. If the heat produced by the action of the oil of vitriol and solution is considered insufficient to expel all the carbonic acid from the liquid, the plug is again inserted in the tube and the contents of the flask gently boiled for some seconds. When the apparatus is nearly cold, more air is again drawn through it, and the whole finally weighed. The loss is due to carbonic acid gas (CO_2), from the weight of which that of any carbonate is ascertained by calculation. Carbonates insoluble in water may be attacked by hydrochloric instead of sulphuric acid; granulated mixtures of carbonates and powdered tartaric or citric acids by enclosing the preparation in the inner tube and placing water in the flask, or *vice versa*. The apparatus also may be modified in many ways to suit the requirements, convenience, or practice of the operator.

Process 2.—Carbonates from which carbonic acid gas is evolved by heat may be estimated by the loss they experience on ignition.

Process 3.—Free carbonic acid gas may be absorbed by a solid stick of potash or strong alkaline solution, the loss in volume of the gas or mixture of gases indicating the amount originally present.

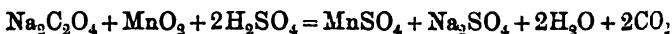
Weights of equivalent quantities of carbonic acid gas and certain carbonates.

Carbonic acid gas	CO_2	44
Carbonic acid	H_2CO_3	62
Anhydrous carbonate of sodium	Na_2CO_3	106
Crystalline carbonate of sodium	$\text{Na}_2\text{CO}_3 \cdot 10\text{H}_2\text{O}$	286
Anhydrous carbonate of potassium	K_2CO_3	138
Crystalline carbonate of potassium	$\text{K}_2\text{CO}_3 + 16\% \text{ Aq}$	164.235
Carbonate of calcium	CaCO_3	100

OXALATES.

Process 1.—The oxalic radical is usually precipitated in the form of oxalate of calcium, and weighed as carbonate, the manipulations being identical with those observed in the estimation of calcium (*vide* p. 681). The experiment may be performed on 0.3 or 0.4 of pure crystallized oxalic acid, 126 parts of which should yield 100 of carbonate of calcium.

Process 2.—Oxalates may also be determined by conversion of their acidulous radical into carbonic acid gas, and observation of the weight of the latter. The oxalate, water, and excess of black oxide of manganese are placed in the carbonic-acid apparatus (p. 702), a tube containing oil of vitriol lowered into the flask, the whole weighed, and the operation completed as for carbonates. From the following equation it will be seen that every 88 parts of carbonic acid gas evolved indicates the presence of 126 parts of crystallized oxalic acid or an equivalent quantity of other oxalate :—



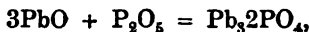
The black oxide of manganese used in this experiment must be free from carbonates. The amount of materials employed is regulated by the size of the vessels.

PHOSPHATES.

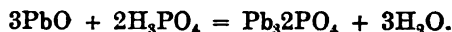
Process 1.—From phosphates dissolved in water the phosphoric radical may be precipitated and weighed in the form of pyrophosphate of magnesium, the details of manipulation being similar to those observed in estimating magnesium (*vide* p. 683). Half a gramme or rather more of pure dry crystallized phosphate of sodium may be employed in experimental determinations. The official phosphate of ammonium (*Ammonice Phosphas*, B. P.) is quantitatively analysed by this method. "If twenty grains of this salt be dissolved in water, and solution of ammonio-sulphate of magnesium added, a crystalline precipitate falls, which, when well washed upon a filter with solution of ammonia diluted with an equal volume of water, dried, and heated to redness, leaves 16·8 grains." Half a gramme, or less, is a more convenient quantity if the operations be conducted with care. Solution of ammonio-sulphate of magnesium (B. P.) is prepared by dissolving 2 parts of sulphate of magnesium, 1 of chloride of ammonium, and 1 of solution of ammonia (20·6 per cent. NH_4HO) in 18 or 20 of distilled water; such a solution is of considerable use if several phosphoric determinations are about to be made.

Process 2.—Free phosphoric acid is most readily determined as phosphate of lead ($\text{Pb}_3\text{2PO}_4$). Of the official solution of phosphoric acid it is stated that "355 grains by weight poured upon 180 grains of oxide of lead in fine powder leave, by evaporation, a residue (principally phosphate of lead) which after it has been heated to dull redness, weighs 215·5 grains." One-tenth of these quantities may be used for experimental purposes; one to two grammes will give good results. The oxide of lead must be quite pure; it should be prepared by digesting red lead in warm dilute nitric acid, washing, drying, and heating the resulting puce-coloured plumbic oxide in a covered porcelain crucible. The increase in weight obtained on evaporating a given amount of solution of phosphoric acid with a known weight of perfectly pure oxide of lead (PbO)

may be regarded as entirely due to phosphoric anhydride (P_2O_5),



the actual reaction being



From these equations, and the table of atomic weights (vide Appendix), the percentage of phosphoric acid (H_3PO_4) in any specimen of its solution may easily be calculated.

Process 3.—The strength of pure solution of phosphoric acid may be ascertained by specific gravity and reference to Tables.

Process 4.—Bone-earth, “superphosphate,” the official *Calcii Phosphas*, and other forms of phosphate of calcium known to be tolerably free from iron or aluminium, may be estimated by treating about half a gramme with hydrochloric acid somewhat diluted, filtering if necessary, warming, precipitating with excess of ammonia, collecting the precipitate ($Ca_3(PO_4)_2$), washing, drying, igniting, and weighing. The phosphate of calcium of pharmacy, if pure, will in this process lose no weight.

Process 5.—Insoluble phosphates in ashes, manures, etc., are treated as follows. The weighed quantity of the material (1.0 to 10.0) is digested in hydrochloric acid diluted with three or four times its bulk of water, filtered (insoluble matter and filter being thoroughly exhausted by water), ammonia added to the filtrate and washings, until, after stirring, a faint cloudy precipitate is perceptible, solution of oxalic acid dropped in until, after agitation for a few minutes, the opalescence is destroyed, oxalate of ammonium next added, the whole warmed, oxalate of calcium removed by filtration, and the filtrate concentrated if very dilute, the liquid treated with citric acid in such quantity that ammonia when added in excess gives a clear lemon-yellow solution (Warington), magnesian mixture poured in (as in process 1), and the precipitate of ammonio-magnesian phosphate collected, washed, dried, and

weighed, as already described in connection with the estimation of magnesium.

Relative weights of equivalent quantities of phosphoric compounds.

Phosphoric acid	H_3PO_4	98
Pyrophosphate of magnesium	$(\text{Mg}_2\text{P}_2\text{O}_7 = 222) \div 2 =$	111
Phosphate of lead	$(\text{Pb}_3\text{P}_2\text{O}_4 = 811) \div 2 =$	405.5
Phosphoric anhydride . . .	$(\text{P}_2\text{O}_5 = 142) \div 2 =$	71
Phosphate of calcium . . .	$(\text{Ca}_3\text{P}_2\text{O}_4 = 310) \div 2 =$	155
Superphosphate of calcium.	$(\text{CaH}_4\text{P}_2\text{O}_4 = 234) \div 2 =$	117

QUESTIONS AND EXERCISES.

1040. What quantity of pure rock-salt is equivalent to 4.2 parts of chloride of silver?—*Ans.* 1.712.

1041. State the percentage of real iodide of potassium contained in a sample of which 8 parts yield 10.9 of iodide of silver.—*Ans.* 96.25.

1042. What is the strength of a solution of hydrocyanic acid 10 parts of which, by weight, yield .9 of cyanide of silver?—*Ans.* 1.81 per cent.

1043. How are nitrates quantitatively estimated?

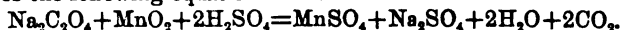
1044. By what processes may the strength of sulphides be determined?

1045. How much real sulphate of sodium is contained in a specimen 10 parts of which yield 14.2 of sulphate of barium?—*Ans.* 86.34 per cent.

1046. Give details of the operations performed in the quantitative analysis of carbonates.

1047. What amount of carbonic acid gas should be obtained from 10 parts of acid carbonate (or bicarbonate) of potassium?—*Ans.* 4.4 parts.

1048. To what operation and what proportions of materials does the following equation refer?



1049. Explain the lead process for the estimation of phosphoric acid in the official solution.

1050. State the amount of superphosphate of calcium equivalent to 7.6 parts of pyrophosphate of magnesium.—*Ans.* 8.01 parts.

SILICATES.

Silica (SiO_2) may be separated from alkaline silicates, or from silicates decomposable by hydrochloric acid, by digesting the substance in hydrochloric acid at a temperature of 70° or 80° C. until completely disintegrated, evaporating to dryness, heating in an air-bath, again moistening with acid, diluting with hot water, filtering, washing, drying, igniting, and weighing.

ESTIMATION OF WATER.

Water and other matters readily volatilized are most usually estimated by the loss in weight which a substance undergoes on being heated to a proper temperature. Thus, in the British Pharmacopœia, crystalline gallic acid ($\text{H}_3\text{C}_7\text{H}_3\text{O}_6$, H_2O) is stated to lose 9.5 per cent. of its weight at a temperature of 100° C., oxalate of cerium (CeC_2O_4 , $3\text{H}_2\text{O}$) 52 per cent. on incineration, carbonate of potassium about 16 per cent. on exposure to a red heat, sulphate of quinine ($2\text{C}_{20}\text{H}_{24}\text{N}_2\text{O}_7$, H_2SO_4 , $7\text{H}_2\text{O}$) 14.4 per cent. at 100° C., arseniate of sodium (Na_2HASO_4 , $7\text{H}_2\text{O}$) 40.38 per cent. at 149° C., carbonate of sodium (Na_2CO_3 , $10\text{H}_2\text{O}$) 60.3 per cent., phosphate of sodium (Na_2HPO_4 , $12\text{H}_2\text{O}$) 63 per cent., and sulphate of sodium (Na_2SO_4 , $10\text{H}_2\text{O}$) 55.9 per cent. at a low red heat: oxide of bismuth heated to incipient redness is not to diminish in weight.

Process.—One or two grammes of substance is sufficient in experiments on desiccation, the material being placed in a watch-glass, covered or uncovered porcelain crucible, or other vessel, according to the temperature to which it is to be exposed.

Rapid desiccation at an exact temperature may be effected by introducing the substance into a tube having somewhat the shape of the letter U, sinking the lower part of the tube into a liquid kept at a definite temperature by aid of a thermometer, and drawing or forcing a current of dry air slowly through the apparatus. Substances liable to oxidation may be desiccated in a current of dried carbonic acid gas. The

weights of the U-tube before and after the introduction of the salt, and after desiccation, give the amount of water sought. In all cases the material must be heated until it ceases to lose weight. Occasionally it is desirable to estimate water directly by conveying its vapour in a current of air through a weighed tube containing chloride of calcium and re-weighing the tube at the close of the operation; the increase shows the amount of water.

Note.—Highly dried substances rapidly absorb moisture from the air; they must therefore be weighed quickly, enclosed, if possible, in tubes (p. 675), a pair of clamped watch-glasses, or a crucible having a tightly fitting lid.

CARBON, HYDROGEN, OXYGEN, NITROGEN.

The quantitative analysis of animal and vegetable substances is either *proximate* or *ultimate*. Proximate quantitative analysis includes the estimation of water, oil, albumen, starch, cellulose, gum, resin, alkaloids, acids, glucosides, ash. It requires the application of much theoretical knowledge and manipulative skill, and cannot well be studied except under the guidance of a tutor. One of the best of the published works on the subject is by Rochleder, a translation of whose monographs will be found in the *Pharmaceutical Journal*, vol. i. 2nd ser. pp. 562, 610; vol. ii. 2nd ser. pp. 24, 129, 160, 215, 274, 420, 478. Another is a book by Professor A. B. Prescott, "Outlines of Proximate Organic Analysis" (Van Nostrand, New York).

Ultimate quantitative organic analysis can only be successfully accomplished with the appliances of a well-appointed laboratory—a good balance, a gas-furnace (p. 691) giving a smokeless flame (7 or 8 centimetres wide and 70 or 80 centims. long), special forms of glass apparatus, etc. *The theory of the operation* is simple: a weighed quantity of a substance is burnt to carbonic acid gas ($\text{CO}_2 = 44$) and water ($\text{H}_2\text{O} = 18$), and these products collected and weighed; 12 parts in every 44 of carbonic acid gas ($= \frac{3}{11}$) are carbon, 2 in every 18 of water ($= \frac{1}{9}$) are hydrogen; nitrogen, if present, escapes as gas. If nitrogen be a constituent, more of the substance is strongly heated with a mixture of the hydrates of sodium and

calcium; these bodies then split up into oxides, oxygen, and hydrogen; the oxygen burns the carbon of the substance to carbonic acid gas, its hydrogen and nitrogen appearing as water and ammonia respectively; the carbonic acid and water are disregarded, the ammonia collected and weighed in the form of a double chloride of platinum and ammonium ($\text{PtCl}_4 \cdot 2\text{NH}_4\text{Cl} = 447$), of which 28 parts in every 447 ($= \frac{1}{16}$) are nitrogen. The difference between the sum of the weights of hydrogen and carbon, and the weight of substance taken, is the proportion of oxygen in the body, supposing nitrogen to be absent. If nitrogen is present, the difference between the sum of the percentages of carbon, hydrogen, and nitrogen and 100 is the percentage of oxygen. Shortly, carbon is estimated in the form of carbonic acid gas, hydrogen as water, nitrogen as ammonia, and oxygen by loss.

The following is the outline of the necessary manipulations. The source of the oxygen for the combustion of carbon and hydrogen is black oxide of copper in coarse powder. 200 or 300 grammes of this material are heated in a crucible to low redness for a short time to expel every trace of moisture—then transferred to store-tubes (fig. 84) resembling test-tubes.

Fig. 84.

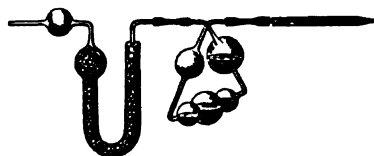
half a mètre long, and having a slightly narrowed mouth, the tube being held in a cloth to protect the hand while the hot oxide is being directly introduced into the mouth of the tube by a scooping motion. As soon as the well-corked tube is cool, the oxide is poured, portion by portion, into a similar tube (the combustion-tube), but somewhat longer, drawn out to a quill (bent upwards nearly to a right angle) at one end, and not constricted at the mouth. Two such tubes are readily made by softening in the blowpipe-flame two or three centimètres of the central part of a tube about a mètre long, and drawing the halves of the tube apart as shown in the following engraving (fig. 85). The tubes are separated by melting the glass in the middle of the quilled portion. A few décigrammes of fused chlorate of potassium should first be dropped into the tube. After ten or fifteen centimètres of

Fig. 85.



oxide have been poured in, about a décigramme of the substance to be analysed is dropped down the tube, then a few grammes of oxide, then another décigramme of substance, then more oxide, until three or four décigrammes of the body under examination have been added. The fifteen or twenty centimètres of alternate layers are next thoroughly mixed by a long copper wire having a short helix; more oxide is introduced, the wire cleansed by twisting the helix about in the pure oxide, and a plug of dried asbestos finally placed on the top of the oxide at about five centimètres from the mouth of the tube; the tube is then securely corked and set aside. The substance operated on may be pure white sugar, powdered and dried; the tube in which it is contained is weighed before and after the removal of a portion for combustion; the loss is the quantity employed in the experiment. The combustion-furnace may be such as shown on page 691. If the furnace is very powerful, or the combustion-tube not of the hardest glass, the tube should be enclosed in wire-gauze the elasticity of which has been destroyed by heating to redness. If the substance under experiment contain nitrogen, the plug of asbestos must be displaced by one of copper turnings, which serves to reduce any oxides of nitrogen, and thus insure the escape of nitrogen itself—or dry yellow chromate of potassium may be used (Perkin). The *water* produced

Fig. 86.



CHLORIDE-OF-CALCIUM TUBE AND POTASH-BULBS.

when the prepared tube is heated is collected in a small U-tube containing pieces of chloride of calcium, or pumice-stone moistened with sulphuric acid (fig. 86); the *carbonic acid gas* in a series of bulbs (fig. 86) containing solution of potash (sp. gr. about 1.27). These bulbs may be purchased at any apparatus-shop. The chloride-of-calcium tube is fitted by a good cork to the combustion-tube, the potash-bulbs by a short piece of india-rubber tubing to the chloride-of-calcium tube. The potash-bulbs may carry a short light tube containing a rod of caustic potash three or four centimètres long; this serves to arrest any moisture that might be carried away from the solution of potash by the dried expanded air which escapes during the operation. The combustion-tube having been placed in the furnace, and the drying-tube and potash-bulbs weighed and attached, the gas is lit under the asbestos, and, when the tube is red-hot, the flame slowly extended until nearly the whole tube is at the same temperature, the operation being conducted at such a rate that bubbles of gas escape through the potash-bulbs at about the rate of one per second. When no more gas passes, the extremity of the tube containing the chlorate of potassium is gently heated until oxygen ceases to be evolved; the quilled extremity of the combustion-tube is then broken, and air drawn slowly through the apparatus by suction through an india-rubber tube fixed on the free end of the potash-bulbs; perfect combustion of carbon and removal of all carbonic acid gas is thus insured. The drying-tube and bulbs are disconnected and weighed, the increase in weight due to carbonic acid gas and water respectively noted, and the percentages of carbon, hydrogen, and (by loss) oxygen calculated. This method is that of Liebig, with modifications by Bunsen; good combustion-furnaces are those known as Hofmann's and Griffin's.

Chromate of lead can be used for combustions in place of oxide of copper. Its advantages are its less hygroscopic nature, and the greater readiness with which it yields its oxygen to organic bodies when heated with them. It must not, however, be used with bodies containing nitrogen, since it would

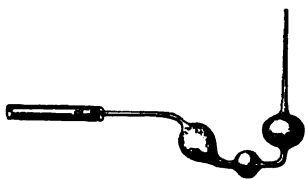
convert so large a proportion of the nitrogen into nitric oxide or higher oxide of nitrogen, that it would be necessary to use an inconveniently long layer of copper turnings to reduce these oxides and so prevent their absorption in the series of bulbs containing the solution of potash. Organic bodies, however, containing sulphur, bromine, iodine, or chlorine are burnt with advantage by means of chromate of lead. If oxide of copper were used with bodies containing sulphur, it would be necessary to place an additional tube containing peroxide of lead between the chloride-of-calcium tube and the potash-bulbs in order to absorb the sulphurous anhydride formed; this is entirely obviated by using chromate of lead, which itself retains the whole of the sulphur. Again, if bodies containing chlorine, iodine, or bromine are burnt by means of oxide of copper, then volatile chloride, iodide, or bromide of copper is formed, and collecting in the chloride-of-calcium tube vitiates the result with regard to the hydrogen; by using chromate of lead, however, the chlorine, iodine, and bromine are respectively retained in the combustion-tube as chloride, bromide, and iodide of lead.

In order to render the chromate fit for use, it is first fused and poured out on a clean iron plate; when cool it is powdered, and heated in a long tube throughout its whole length, while air, dried by passing through chloride of calcium or strong sulphuric acid, is drawn over it; when the colour of the chromate changes to brown the heat can be withdrawn and the extremity of the tube farthest from the drying-apparatus closed, so that the air in passing into the tube on cooling may be quite dry; when cool the drying-tube is removed, the extremity securely corked, and the chromate of lead is ready for direct transference to the combustion-tube.

The general manipulations for substances containing nitrogen resemble the foregoing so far as the use of a combustion-tube and furnace and collection of the ammoniacal gas are concerned. The combustion-tube must be quilled at one end, and about a third of a mètre long. The soda-lime is made by slaking quicklime with a solution of soda, of such a strength

that about two parts of quicklime shall be mixed with one of hydrate of sodium, drying the product, heating to bright redness, and finely powdering; it should be preserved in a well-closed bottle. Some of the soda-lime is introduced into the tube, then layers of substance and soda-lime, mixture effected

Fig. 87.



NITROGEN-BULBS.

by a wire, more soda-lime added, and lastly a plug of asbestos. Bulbs (fig. 87), known as those of Will and Varrentrapp (the originators of the method), containing hydrochloric acid of about 25 per cent., are then fitted by a cork, and the tube heated in the furnace. When

gas ceases to pass, the quill is broken, and aspiration continued slowly until all ammoniacal gas may be considered to have been absorbed by the acid. The bulbs are disconnected, their contents and rinsings poured into a small dish, solution of perchloride of platinum added, and the operation completed as in the estimation of ammonium and potassium salts (*vide* pp. 679 and 676). (See a paper "On the determination of nitrogen," in the *Chemical News* of July 18, 1879.)

Liquids are analysed by a similar method to that adopted for solids, volatile liquids being enclosed in small bulbs having a long quill. These are weighed previously to and after the introduction of the liquid; just before being dropped into the combustion-tube the quill is broken.

Formulae.—From the percentage composition of an organic substance an empirical formula may be deduced by dividing the weight of each constituent by its atomic weight, and converting the product into the simplest whole numbers; a rational formula by ascertaining the proportion in which the substance unites with a radical or body having a known combining proportion, etc. (*vide* p. 504).

Chlorine, bromine, or iodine contained in an organic substance is usually estimated by heating to redness a given weight of the material with ten times as much pure lime in a

combustion-tube. Chloride, bromide, or iodide of calcium is thus produced. While still hot the tube is plunged into water, the mixture of broken glass and powder treated with pure diluted nitric acid in very slight excess; the filtered liquid precipitated by nitrate of silver, and the chloride, bromide, or iodide of silver collected, washed, dried, and weighed.

Sulphur, phosphorus, and arsenicum in organic salts may be estimated by gradually heating in a combustion-tube 1 part of the substance with a mixture of 10 parts nitre, 2 dried carbonate of sodium (in order to moderate deflagration), and 30 chloride of sodium. The product is dissolved in water acidulated by nitric acid, the sulphuric radical precipitated and estimated as sulphate of barium, the phosphoric and arsenic radicals as ammonio-magnesian phosphate or arseniate.

QUINIA OR QUININE.

De Vrij's Method for the Separation of the Mixed Alkaloids from Cinchona Barks.—Twenty grammes of powdered and sifted bark, dried at 100° C., are mixed with milk of lime, made of 5 grammes of dry slaked lime and 50 grammes of water. This mixture is slowly dried, and, when entirely dry, heated in a flask with 200 c. c. of very strong alcohol (about 95 per cent.) till it boils. After well shaking, the clear liquid is poured by degrees into a percolating tube. The flask is then washed with 200 c. c. more of the same alcohol, and the liquid passed through the percolator. After this has run through, a little clear colourless liquid may be further collected by pouring into the percolator successive small quantities of distilled water. The united clear liquors are now slightly acidulated with weak sulphuric acid, whereby a precipitate of sulphate of calcium is formed. When this has subsided, the greater part of the liquid may be decanted and the rest passed through a small filter. From the clear solution most of the spirit is distilled off, the residue poured into a capsule, and the still washed with a little weak spirit, the washings being added to the liquid in the capsule. The capsule is now heated on a water-bath till free from alcohol, and the remaining liquor, which contains all the alkaloids in the form of acid sulphates, is, after cooling, passed through a small filter. On the filter

remains a mixture of quinovine* and fatty substances, which must be washed with water slightly acidulated with sulphuric acid, till the washings are not rendered turbid by caustic soda. The mixed filtrate and washings, concentrated over a water-bath, is, while *still warm*, precipitated by a large excess of caustic soda. This precipitation from a warm solution has the advantage of rendering the alkaloids less voluminous, and therefore capable of being more easily washed. (Should the alkaloids melt—as is sometimes the case under this treatment—the liquid must be cooled and the fused alkaloids carefully powdered before proceeding to the next step of the process.) The precipitate is next placed on a small filter, washed with the smallest possible quantity of distilled water (sufficient to remove the soda salt without dissolving the quinine, etc.): the filter is laid on blotting-paper, and this is renewed until the mixed alkaloids can easily be removed from the filter—an operation that can be accomplished before they are quite dry, but requires some practice. They are then heated in a tared capsule on a water-bath till they cease to lose weight. The observed weight multiplied by five gives the percentage of mixed alkaloids in the bark. The intensely yellow alkaline liquor separated from the alkaloids contains still a small quantity of quinovine, which can be separated by acidifying with hydrochloric acid.

De Vrij's Method for the Separation and Quantitative Determination of all the different Cinchona Alkaloids.—This is based upon the following facts (nomenclature adopted by the Quinological Congress of Amsterdam, 1877):—

1. The great solubility of quinine and amorphous alkaloids in ether, and the *relative* insolubility of quinidine, cinchonidine, and cinchonine in this liquid.

2. The great solubility of the iodosulphate of the amorphous alkaloid in alcohol of at least 90 per cent., and the very slight solubility of the iodosulphate of quinine (herapathite) in the same.

3. The great difference in solubility between the tartrate of cinchonidine and the tartrates of cinchonine and quinidine—the first being soluble in 1265 parts of water at 10° C., the second in 35.6 parts of water at 16° C., and the third in 38.8 parts of water at 15° C.

* "This is strictly a mixture of quinovine and quinovic acid, but as quinovine is by far the larger constituent, I prefer the name of quinovine for the mixture."—*De Vrij*.

4. The great difference in solubility between the hydriodate of quinidine and the hydriodates of cinchonidine and cinchonine in water and alcohol. One part of hydriodate of quinidine requires 1250 parts of water at 15° C. or 110 parts of alcohol; the same salt of cinchonidine 110 parts of water or 3 parts of alcohol; and that of cinchonine 128 of water or 3 of alcohol.

These facts are applied to the separation and determination of the different cinchona alkaloids in the following manner:—

(A.) *Determination of Quinine.*—Two grammes of the pulverized mixed alkaloids are mixed with 30 grammes of ether, and the mixture, after well shaking, left at rest till the next day. By this operation all the quinine and amorphous alkaloid are dissolved, together with traces of quinidine and cinchonidine, while the chief part of these, together with cinchonine, remains undissolved.

After filtering the ethereal solution and washing the residue with ether, the united filtrates are distilled and the weight of the residue ascertained. This is dissolved in 40 times its weight of alcohol of 90 or 92 per cent., containing .7562 per cent. of sulphuric acid, to obtain an alcoholic solution of the acid sulphates of the alkaloids. From this solution the quinine is separated by adding carefully by means of a pipette, an alcoholic solution of *iodosulphate of chiniodine*,* as long as

* This reagent is thus prepared: Two parts of *Sulphate of Chiniodine* are dissolved in 8 parts of water, containing 5 per cent. of sulphuric acid. (The salt is identical with *Sulphate of Amorphous Quinine*, prepared according to Dr. de Vrij's process, by Messrs. Howard and Sons, Stratford.) To this clear solution, contained in a large capsule, a solution of one part of iodine and two parts of iodide of potassium in 100 parts of water is slowly added under continuous stirring so that no part of the solution of chiniodine comes into contact with an excess of iodine. By this addition an orange-coloured flocculent precipitate is formed of iodosulphate of chiniodine, which either spontaneously or by a slight elevation of temperature collapses into a dark brown-red coloured resinous substance, whilst the supernatant liquor becomes clear and slightly yellow-coloured. This liquor (still containing chiniodine, which can be obtained in a very pure state, if a little sulphurous acid is added before precipitating the alkaloid by caustic soda) is poured off, and the resinous substance is washed by heating it on a water-bath with distilled water. After washing, the resinous substance is heated on the water-bath till all the water has been evaporated. It is then soft and tenacious at the temperature of the water-bath, but becomes hard and brittle after cooling. One part of this substance is now heated with 6 parts of alcohol of 92 or 94 per cent. on a water-bath, and is thus dissolved, and the solution allowed to cool. In cooling, a part of the dissolved substance is separated. The clear dark-coloured solution is evapor-

a dark brown or red precipitate of iodosulphate of quinine (herapathite) falls. As soon as all the quinine has been precipitated and a slight excess of the reagent added, the liquid acquires an intense yellow colour. The beaker containing the liquor and precipitate is now covered with a watch-glass and heated on a water-bath till the liquid *begins* to boil. After cooling the beaker is weighed to ascertain the amount of liquid present; the latter is then poured on to a filter placed in a tared funnel, and the precipitate washed with a saturated solution of herapathite in alcohol (92 per cent.). The weight of the moist filter is now taken, and the precipitate allowed to dry in the funnel, and its weight again taken. It is then removed from the funnel to a watch-glass and completely dried over a water-bath; another closely fitting glass is then placed over it and its weight is once more taken, the weighing being repeated till constant. The *rationale* of the latter part of the process is as follows: Herapathite is soluble in alcohol at 15° C. to the extent of .125 per cent. The quantity of herapathite held in solution by the alcohol present after the addition of the precipitant used is to be calculated and added to the final amount obtained. The difference in weight between the moist and the dry filter (weighed in the funnel) gives the amount of alcohol driven off in drying, and from this can be calculated the amount of herapathite left on the filter from the saturated solution employed in the washing. A final correction can then be made by subtracting this from the sum obtained as above—the temperature of the laboratory being taken into account in both cases. The resulting correction may be thus expressed:—

$$\left. \begin{array}{l} \text{Total weight} \\ \text{of herapathite} \end{array} \right\} = \text{wt. obtained} + .00125 \times \left\{ \begin{array}{l} \text{total weight of al-} \\ \text{cohol present} - \\ \text{loss of alcohol in} \\ \text{drying precip.} \end{array} \right.$$

From this weight the amount of *crystallisable* quinine can be calculated by the use of Jörgensen's formula— $4\text{C}_{20}\text{H}_{24}\text{N}_2\text{O}_2 \cdot 3\text{H}_2\text{SO}_4, 2\text{HI}, \text{I}_4 = 2354$; according to which 1 part of iodosul-

ated on a water-bath, and the residue dissolved in 6 parts of cold alcohol. This second solution leaves a small part of insoluble substance. The clear dark-coloured solution obtained by the separation of this insoluble matter, either by decantation or filtration, constitutes the reagent which De Vrij, under the name of *iodosulphate of chiniodine*, uses both for the qualitative and quantitative determination of *crystallisable* quinine.

phate of quinine, dried at 100° C., represents '55055 of anhydrous quinine, or '734 of pure commercial disulphate of quinine.

(B.) *Determination of the other Alkaloids.*—Two grammes of the pulverized mixed alkaloids are dissolved in weak hydrochloric acid to obtain a slightly alkaline solution measuring 70 c.c. By adding 1 gramme of Rochelle salt to this solution the tartrates of quinine and cinchonidine are separated; these are collected on a filter, washed with a little water, and dried on a water-bath. One part of these tartrates represents '80844 of quinine and cinchonidine: from the amount of these alkaloids thus found the amount of quinine already ascertained is subtracted, the remainder representing the cinchonidine present. In the filtrate from the tartrates quinidine (if present) is precipitated by a concentrated solution of potassium iodide; one part of the dried hydriodate representing '86504 of crystallized quinidine. The remaining solution is treated with caustic soda, and the precipitate (if any) washed with ether. The residue represents the amount of cinchonine. Finally, by distilling the ether from the washings can be ascertained the amount of amorphous alkaloid, which often, in the case of analysis of Indian barks, contains traces of quinamine.

Carles's process for the valuation of Cinchona Barks.—An average sample of the bark is reduced to fine powder and passed through a sieve without residue. Twenty grammes are then taken and intimately mixed in a mortar with 8 grammes of slaked lime previously mixed with 35 grammes of water. This mixture spread on a plate is dried in the air in summer, or on a water-bath at other times. When all the moisture has evaporated the lumps are broken up and the powder packed in a percolator with a piece of lint at the bottom. Chloroform is then passed through in successive portions till the mass is exhausted. This is ascertained by receiving the last drops on a watch-glass, evaporating to dryness and pouring on the residue water acidulated with dilute sulphuric acid, then solution of chlorine, and lastly ammonia. When a green colour is no longer produced it is known that all the quinia has been removed. When the operation is well conducted about 150 grammes of chloroform suffice for this purpose. The menstruum retained by the mass is displaced by water, and the whole of the chloroform solution is either distilled or evaporated to dryness. To separate the

alkaloids from the residue it is treated several times in the cold with diluted sulphuric acid (1 to 10) 10 to 12 cubic centimètres being sufficient. This solution thrown upon a moistened filter passes through colourless and free from resinous matter. It is raised to the boiling point and ammonia cautiously added so as to leave the liquid with a *slightly acid* reaction. The sulphate of ammonium thus formed appears to prevent the mother-liquors from retaining sulphate of quinine in solution. All the quinine crystallizes out in the state of sulphate. After some time it is collected on a double filter, the mother-liquors displaced by a little water, and the crystals dried and weighed. It is preferable to dry completely at 100° C., and after weighing in this state to add the 12 per cent. of water, which is lost by this treatment. The other alkaloids retained in the mother-liquors are separated by precipitation.

A quantitative determination of the purity of commercial Sulphate of Quinine may be made by dissolving 1 gramme of the salt in 20 grammes of alcohol containing .7562 per cent. of sulphuric acid, adding 20 more grammes of alcohol, and precipitating the quinine as herapathite in the manner above described. If pure, one gramme of quinine should yield 1.3 gramme of herapathite (De Vrij).

Kerner's Test for the purity of Sulphate of Quinine.—This has already been described as a qualitative test under quinine (p. 457). It may be applied quantitatively in the following manner: 5 grammes of the sulphate to be tested is triturated in a mortar with a little cold water, and the whole placed in a stoppered bottle previously calibrated for 50 c. c. Water is added to 50 c. c. and the mixture is well agitated and set aside for a day or so. It is then filtered and a portion of the filtrate, say 30 c. c., is examined. To 30 c. c. of the filtrate 42 c. c. of ammonia solution (sp. gr. 0.96) is added, and the mixture agitated. If a permanent precipitate remains it is collected on a filter, washed, dried at 100° C., and weighed. This weight is the amount of alkaloid other than quinine in 3 grammes of sulphate. Multiplied by 33.3 it is the percentage of alkaloid other than quinine, generally cinchonidine, contained in the specimen of sulphate. To obtain the amount of sulphate of cinchonidine corresponding to this percentage it must be multiplied by 1.217.

Paul's Modification of Kerner's Test for Sulphate of Cinchonidine in Sulphate of Quinine.—Dissolve 4 or 5 grammes of

the salt in 80 to 150 c. c. of boiling water. Set aside for some hours, or during a night. Most of the quinine sulphate is deposited, while the more soluble cinchonidine sulphate remains in solution. Filter. To the filtrate in a bottle add ether, shaking occasionally, until a distinct layer of ether remains undissolved. Add now ammonia in excess; quinine is precipitated and then redissolved by the excess of ammonia, while cinchonidine remains insoluble. Set this mixture aside for a few hours; any cinchonidine which may have been carried into solution in the ether along with the quinine will separate out in a crystalline state. If a weighed quantity of the original salt has been operated on, the separate cinchonidine may be collected on a filter, washed with a little ether, and weighed. Care must be taken that the filter does not retain quinine deposited by the evaporation of the ether which held it in solution. This will be avoided if the flask in which the operation is conducted is of such a size that the ethereal solution of quinine collects in the neck; for the latter liquid may be removed by a pipette, a little more ether then poured in, the cork replaced, and the whole shaken and set aside; and this operation repeated two or three times before the cinchonidine is collected. The cinchonidine being thus washed with a minimum of ether suffers a minimum loss.

The quinine sulphate deposited from the first aqueous solution should be again recrystallized, and its mother-liquor treated as above. And even the sulphate deposited in this second treatment should be once more carried through the operation. The whole of the cinchonidine may finally be collected on one filter.

The aqueous filtrates may even be evaporated, set aside, filtered, and the ether test again applied (Parker).

Sulphate of quinine normally contains 14.45 per cent. of water; sulphate of cinchonidine 13.17 per cent.; all given off at 100° to 115° C. The drying should therefore be effected at 100° C., and the dried salt weighed in well-fitting weighing tubes. 100 parts of cinchonidine are equivalent to 116 parts of sulphate of cinchonidine.

Sulphate of cinchonidine is almost the only salt likely to be accidentally present in sulphate of quinine, much quinine being rarely present in bark, and sulphate of cinchonine being sufficiently soluble to always remain in the mother-liquors of sulphate of quinine.

Of the Citrate of Iron and Quinia (Ferri et Quinice Citrus, B. P and U. S. P.) it is officially stated that "fifty grains dissolved in a fluid ounce of water and treated with a slight excess of ammonia give a white precipitate which, when collected on a filter, washed, and dried (at 260° F.), weighs eight grains. The precipitate is almost entirely soluble in two or three fluid drachms of pure ether," and the ethereal solution set aside for twelve hours in a small well-corked bottle yields no crystalline deposit (of quinidia).

Another process for the determination of the quinia of the scaled compound. A weighed quantity of the scale, about 4 grammes, is dissolved in about 50 c. c. of water, and the whole is placed in a closed separating funnel. About the same volume of chloroform is added and enough ammonia to impart a distinctly alkaline reaction. The whole is well agitated and is allowed to stand until the two layers separate. The chloroformic layer is then run into a weighed dish. The aqueous solution is treated in this way with two more portions of chloroform about 25 c. c. each. The mixed chloroformic solutions are then evaporated to dryness over a water-bath, and the weight of the residue determined. To the residue is added about 25 c. c. of water and enough dilute sulphuric acid to impart a decidedly acid reaction. The mixture is next heated over a water-bath until, the solution remaining acid, the residue has completely dissolved. Dilute soda solution is afterwards added with great care until the solution is exactly neutral. The dish is then removed and the solution allowed to cool and rest over night, when the quinine will have separated in crystals of ordinary sulphate. These should be collected on a filter, and the mother-liquor tested with litmus paper. If it is acid it must be warmed over a water-bath and dilute soda solution added to exact neutralization and the solution set aside as before, when some more crystals will probably separate. These are also collected and with the former ones washed, dried at 100° C. and weighed [$(C_{20}H_{24}N_2O_3)_2, H_2SO_4 = 746$]. To this weight must be added 1 gramme for every 750 c.c. of mother-liquor for quinia sulphate which it retains. From this weight of anhydrous quinia sulphate is calculated its equivalent of hydrous quinia $(C_{20}H_{24}N_2O_3)_2, 2H_2O = 684$, the approximate formula of hydrous quinia dried over a water-bath. The weight thus obtained is compared with the weight of total alkaloid determined, both having been reduced to percentages. The amount of hydrous quinia calculated from

the crystals of sulphate should not be much below that weighed directly. In good specimens the difference will be about one per cent.

(See also a paper by Fletcher in the *Pharmaceutical Journal* for Sept. 20, 1879.)

MORPHIA.

The official process for the estimation of this alkaloid in opium is conducted in the following manner:—

Take of opium 100 grains, slaked lime 100 grains, distilled water 4 ounces. Break down the opium, and steep it in an ounce of the water for twenty-four hours, stirring the mixture frequently. Transfer it to a displacement-apparatus, and pour on the remainder of the water in successive portions, so as to exhaust the opium by percolation. To the infusion thus obtained, placed in a flask, add the lime, boil for ten minutes, filter. Wash the residue on the filter with an ounce of boiling water. Acidulate the mixed filtrates slightly with diluted hydrochloric acid, evaporate it to the bulk of half an ounce, and let it cool. Neutralize cautiously with solution of ammonia, carefully avoiding an excess; remove by filtration the brown matter which separates, wash it with an ounce of hot water, mix the washings with the filtrate, concentrate the whole to the bulk of half an ounce, and add now solution of ammonia in slight excess. After twenty-four hours collect the precipitated morphia on a weighed filter, wash it with cold water, and dry it at 212° F. It ought to weigh at least from six to eight grains.

Of Hydrochlorate of Morphia it is stated that "twenty grains of the salt, dissolved in half an ounce of warm water, with ammonia added in the slightest possible excess, give on cooling a crystalline precipitate which, when washed with a little cold water and dried by exposure to the air, weighs 15·18 grains."

Prollius's Method for the Determination of Morphia in Opium.—5 to 10 grammes of opium are weighed and extracted with 100 c. c. of 34 per cent. spirit. The resulting tincture is well shaken with 50 c. c. of ether and 2 c. c. of solution of ammonia in a stoppered bottle and then allowed to stand from twelve to twenty-four hours. The liquids separate slowly and retain, partly in the ether and partly in the alcoholic liquid, the colouring matter, narcotine and other crystallizable constituents of opium, while the morphia separates in crystals

between the two layers and finally sinks to the bottom. The fluid portion is decanted, the crystals are washed with diluted alcohol dried at 100 C., and weighed.

Petit's Method.—Take 15 grammes of opium, suspend it in 75 grammes of water, and afterwards throw it upon a filter. Take 55 grammes of the filtrate, which represents 10 grammes of opium, add 3 c. c. of ammonia solution and agitate. The morphia deposits rapidly in the form of a crystalline powder. The whole is allowed to stand for a quarter of an hour, and then 27 grammes of alcohol (of 95 per cent.) are added; after shaking several times, it is again allowed to stand for half an hour, and then thrown upon a tared filter. The alkaloid is washed upon the filter with 50 per cent. alcohol. It is then dried at 100 C. and weighed.

Flückiger's Method.—Take of powdered opium 8 grammes, cold water 80 grammes; shake the mixture frequently; after twelve hours, filter. The filter should have a diameter of five inches. The operation will afford on an average 65 to 70 grammes of clear liquid. No washing is to take place. 42.5 grammes of the liquid are collected in a little phial, the weight of which has previously been noted. Next add to the solution 12 grammes of alcohol (sp. gr. 0.812–0.815), 10 grammes of ether, and 1.5 gramme ammonia water of 0.960 sp. gr. The mixture after shaking will remain clear and a colourless layer of ether will appear on its surface. The phial is corked and allowed to stand without further shaking. After an hour or two crystals of morphia begin to be formed, mostly at the junction of the two layers. Presently they sink to the bottom, and after a day or two the whole amount of whitish or white crystals of the alkaloid will have been deposited. They are then to be collected by using two folded filters having a diameter of four inches. The phial is rinsed out with a mixture of 6 grammes of alcohol and 5 grammes of ether, and lastly with 10 grammes of ether, these liquids being gradually poured on to the crystals in order to wash them. The funnel in the meantime is carefully covered. The crystals are subsequently cautiously pressed between the folds of the two filters, which will almost completely absorb the mother-liquor which the crystals of morphia may still retain. It will now be quite easy to remove the alkaloid neatly from the filter; it must be weighed in the phial, in which some crystals may have remained obstinately attached to the walls. The phial and contents, dried

at 100° C., contains the whole amount of morphia precipitated, in the hydrous condition, namely, $C_{17}H_{19}NO_3 + H_2O$. The operator finally ascertains the purity of the crystals. To this end, he takes 1 décigramme of the morphia and dissolves it in 10 grammes of lime-water. If the lime-water be duly saturated, in the cold, the quantity mentioned will be a little more than sufficient. The morphia will probably leave only a very trifling amount of colouring matter, quite insufficient to influence appreciably the percentage of alkaloid. Should narcotine be present it will remain undissolved.—It is important to precipitate the morphia from a solution containing alcohol and ether. By adding ammonia to an aqueous solution, a flocculent matter is precipitated. This abundant amorphous mass—an alkaloid or not it is certainly far from being morphia—remains in solution if the liquid contains a little alcohol, one-third alcohol of the volume of the aqueous filtrate being quite sufficient for the purpose. Of no less importance is the action of the ether. It not only prevents the narcotine from being thrown down together with the morphia, but ether promotes the formation of distinct and pure crystals of morphia, this alkaloid separating very readily from a liquid saturated with ether.

SUGAR.

The qualitative test for sugar, by means of an alkaline copper solution (*vide* p. 484), may be applied in the estimation of sugar in sacchariferous substances.

Process 1.—34·65 grammes of pure dry crystals of ordinary sulphate of copper are dissolved in about 250 c. c. of distilled water. 173 grammes of pure crystals of the double tartrate of potassium and sodium are dissolved in 480 c. c. of solution of caustic soda of sp. gr. 1·14. The solutions are mixed and water added to 1 litre. 100 c. c. of this solution represent 3·464 grammes of sulphate of copper, and correspond to 0·5 of a gramme of pure anhydrous grape-sugar, 0·475 of cane-sugar, 0·82 of maltose, or 0·45 of starch. It must be preserved in a well-stoppered bottle to prevent absorption of carbonic acid, and be kept in a dark place. Should it give a precipitate on boiling, a little solution of soda may be added when making experiments.

Dissolve 0.475 of pure dry powdered cane-sugar in about 50 c. c. of water, convert into grape-sugar by acidulating with sulphuric acid and boiling for an hour or two, make slightly alkaline with carbonate of sodium, and dilute to 100 c. c. Place 10 c. c. of the copper solution in a small flask, dilute with three or four times its bulk of water and gently boil. Into the boiling liquid drop the solution of sugar from a burette, one cubic centimètre, or less, at a time, until, after standing for the precipitate to subside, the supernatant liquid has just lost its blue colour; 10 c. c. of the solution of sugar should be required to produce this effect, = 0.0475 of cane-sugar or 0.05 of grape-sugar. Experiments on pure cane-sugar must be practised until accuracy is attained; syrups, diabetic urine, and saccharated substances containing unknown quantities of sugar may then be analysed.

Starch is converted into grape-sugar by gentle ebullition with dilute acid for eight or ten hours, the solution being finally diluted so that one part of starch, or rather sugar, shall be contained in about 150 of water.

Process 2.—Roberts' Method for the Estimation of Sugar in Urine.—About four ounces of saccharine urine are put into a twelve ounce bottle, and a lump of German yeast about the size of a cob-nut or small walnut is added. This excess of yeast hastens fermentation, and does no harm. The bottle is then covered with a grooved cork (to allow of the escape of carbonic acid gas), and set aside in a warm place to ferment. By the side of it is placed a tightly corked four-ounce phial filled with the same urine without any yeast. In about twenty-four hours the fermentation will have ceased, and the scum cleared off or subsided. The fermented urine is then decanted and its specific gravity taken. At the same time the specific gravity of the unfermented urine in the companion phial is observed. The *density lost* is thus ascertained. Each degree of density lost represents a grain of glucose per fluid ounce.

Sugar is often estimated by the measurement of the carbonic acid gas evolved during fermentation.

Saccharometry.—A generic term for certain volumetric

operations undertaken with the view of ascertaining the quantity of sugar present in any matter in which it may be contained.

Saccharometry is frequently performed upon common syrup (*Syrupus*, B. P.) and solutions which are known to contain nothing but cane- (ordinary) sugar, the object being merely to ascertain the amount present. In such a case it is only necessary to take the specific gravity of the liquid at 60° F., and then refer to a previously prepared Table of densities and percentages.

Specific gravity.	Sugar per cent.	Specific gravity.	Sugar per cent.	Specific gravity.	Sugar per cent.
1·007 .	1·8	1·100 .	23·7	1·210 .	46·2
1·014 .	3·5	1·108 .	25·6	1·221 .	48·1
1·022 .	5·2	1·116 .	27·6	1·231 .	50·0
1·029 .	7·0	1·125 .	29·4	1·242 .	52·1
1·036 .	8·7	1·134 .	31·5	1·252 .	54·1
1·044 .	10·4	1·143 .	33·4	1·261 .	56·0
1·052 .	12·4	1·152 .	35·2	1·275 .	58·0
1·060 .	14·4	1·161 .	37·0	1·286 .	60·1
1·067 .	16·3	1·171 .	38·8	1·298 .	62·2
1·075 .	18·2	1·180 .	40·6	1·309 .	64·4
1·083 .	20·8	1·190 .	42·4	1·321 .	66·6 ?
1·091 .	21·8	1·199 .	44·3	1·330 (B.P.)	66·6 ?

The sp. gr. may be taken by an hydrometer, technically termed a *saccharometer*. (The above spec. gravs. = 1° to 35° Baumé.)

If a liquid contains other substances besides cane-sugar, the test of specific gravity is of little or no value. Advantage may then be taken of the fact that syrup causes right-handed twisting of a ray of plane polarized light to an extent exactly proportionate to the amount of sugar in solution. The saccharine fluid is placed in a long tube having opaque sides and transparent ends; and a ray of homogeneous light, polarized by reflection from a black-glass mirror or otherwise, is sent through the liquid and optically examined by a plate of tourmaline, Nicol's prism, or other polarizing eyepiece. Attached

to the eyepiece is a short arm which traverses a circle divided into degrees. The eyepiece and arm are previously so adjusted that when the ray is no longer visible the arm points to the zero of the scale of degrees. The saccharine solution, however, so twists the ray as to again render it visible; and the number of degrees which the eyepiece has to be rotated before the ray is once more invisible is exactly proportionate to the strength of the solution. The value of the degrees having been ascertained by direct experiment and the results tabulated, a reference to the Table at once indicates the percentage of sugar in the liquid under examination. Grape-sugar also possesses the property of dextral rotation, but less powerfully than cane-sugar; moreover the former variety does not, like cane-sugar, suffer inversion of the direction of rotation on the addition of hydrochloric acid to its solution—an operation that furnishes data for ascertaining the amounts of cane- and of grape-sugar, or of crystallizable and non-crystallizable sugar, present in a mixture. In using the polariscope saccharometer, it is convenient to employ tubes of uniform size, and always to operate at the same temperature.

Various modes are adopted of applying, for the purposes of quantitative analysis, this action of syrup on polarized light.

ALCOHOL.

Mulder's process for the determination of the amount of alcohol in wines, beer, tinctures, and other alcoholic liquids containing vegetable matter is as follows:—Take the specific gravity and temperature of the liquid, and measure off a certain quantity (100 cubic centimètres); evaporate to one half or less, avoiding ebullition in order that particles of the material may not be carried away by the steam. Dilute with water to the original bulk, and take the specific gravity at the same temperature as before. Of the figures representing this latter specific gravity, all over 1.000 show to what extent dissolved solid matter affected the original specific gravity of the liquid. Thus, the specific gravity of a sample of wine at

15.5° C. is 0.9951; evaporated till all alcohol is removed and diluted with water to the original bulk, the specific gravity at 15.5° C. is 1.0081: 0.0081 represents the gravitating effect of dissolved solid matter in 0.9951 part of original wine. 0.0081 subtracted from 0.9951 leaves 0.987, which is the specific gravity of the water and alcohol of the wine. On referring to a Table of the strengths of diluted alcohol of different specific gravities, 0.987 at 15.5° C. is found to indicate a spirit containing 8 per cent. of real alcohol. If the foregoing operation be conducted in a retort, the liquid being boiled and the steam carefully condensed, the distillate, diluted with water to the original bulk of wine operated on, will still more accurately represent the amount of water and alcohol in the wine—its specific gravity showing the percentage of real alcohol present.

DIALYSIS.

Dialysis (from *διὰ*, *dia*, through, and *λύσις*, *lusis*, a loosing or resolving) is a term applied by Graham to a process of analysis by diffusion through a septum. The apparatus used in the process is called a *dialyzer*, and is constructed and employed in the following manner. The most convenient septum is the commercial article known as *parchment paper*, made by immersing unsized paper for a short time in sulphuric acid; it is sold by most dealers in chemical apparatus. A piece of this material is stretched over a gutta-percha hoop, and secured by a second external hoop. Dialyzers of useful size are one or two inches deep and five to ten inches wide. Liquids to be dialyzed are poured into the dialyzer, which is then floated in a flat dish containing distilled water. The portion passing through the septum is termed the *diffusate*, the portion which does not pass through is termed the *dialysate*.

The practical value of dialysis depends upon the fact that certain substances will diffuse through a given septum far more rapidly than others. Uncrystallizable bodies diffuse

very slowly. Of such matters as starch, gum, albumen, and gelatine, the last named is perhaps least diffusive; hence substances of this class are termed *colloids*, or bodies like *collin*, which is the soluble form of gelatine. Substances which diffuse rapidly are mostly crystalline; hence bodies of this class are termed *crystalloids*.

Solutions of two parts of the following named substances in 100 parts of distilled water were dialyzed by Graham for twenty-four hours. The amounts of each substance which passed through the septum bore the following relations to one another.

Chloride of sodium	1000
Ammonia	847
Theine	703
Salicin	503
Cane-sugar	472
Amygdalin	311
Extract of logwood	168
Catechu	159
Extract of cochineal	51
Gallo-tannic acid	30
Extract of litmus	19
Purified caramel	5

Ten per cent. solutions, under similar circumstances, gave the following results:—

Gum Arabic	4
Starch-sugar	266
Cane-sugar	214
Glycerine	440
Alcohol	476
Chloride of sodium	1000

Dialyzed iron, an aqueous solution of about 5 per cent. of highly basic oxychloride of iron, is obtained by saturating solution of perchloride of iron with ferric hydrate, by adding ammonia, or, better, carbonate of sodium, and shaking vigorously until the precipitated hydrate ceases to redissolve,

filtering if necessary, placing on a dialyzer floating in distilled water, and displacing the fluid in the dish by water daily for a week or two, or until the diffusate gives no reaction with nitrate of silver. The crystalloids (chloride of sodium or other salt) pass through the dialyzer; the colloid fluid which does not pass through the dialyzer is the highly basic oxychloride of iron, or so-called "dialyzed iron," or "dialytic iron." This fluid has very little taste of iron. Its value as a medicine has been questioned, its non-diffusibility suggesting that it never passes out of the intestinal canal, and therefore never gets into the blood.

The phenomena of dialysis show that crystalloids are superior to colloids in affinity for water. If a solution of chloride of sodium be placed at the bottom of a jar, and covered by a hot solution of gelatine of sufficient strength to solidify on cooling, the chloride of sodium will diffuse up into the solid jelly, because the water of the solid jelly has a greater affinity for the salt than it has for the gelatine. The solid jelly may obviously be reduced in thickness, and saline liquids placed above it; indeed the conditions would then be still more favourable for diffusion. Replace the stratum of jelly by a permanent colloid, such as parchment paper; the result is the same; but the permanent character of the septum admits of its practical application.

Further researches on dialysis will probably throw much light on several important points in connection with physiological chemistry; for there is little doubt that alimentary matter passes through the cell-walls of animals and plants by this process.

QUESTIONS AND EXERCISES.

1051. Carbonate of potassium is said to lose 16 per cent. of water on exposure to a red heat; give the details of manipulation observed in verifying this statement.

1052. Write a few paragraphs descriptive of the process of ultimate organic analysis.

1053. In what forms are carbon, hydrogen, and nitrogen weighed in quantitative analysis?

1054. In the combustion of .41 of a gramme of sugar, what weights of products will be obtained?—*Ans.* .632 of carbonic acid gas (CO_2) and .237 of water (H_2O).

1055. How is cinchona assayed for mixed alkaloids?

1056. On what facts does De Vrij found his method for the separation and quantitative determination of all the cinchona alkaloids?

1057. Describe De Vrij's process for the assay of commercial sulphate of quinia.

1058. Give the official method for the estimation of morphia in opium.

1059. Mention the operations necessary for the estimation of the proportion of sugar in saccharated carbonate of iron, or in a specimen of diabetic urine.

1060. What is understood by *saccharometry*?

1061. Give two processes for the estimation of the percentage of alcohol in tinctures, wines, or beer.

1062. Define dialysis.

CONCLUSION.

Detailed instructions for the quantitative analysis of potable water, articles of food, general technical products, special minerals, soils, manures, air, illuminating agents (including solid fats, oils, spirits, petroleum, and gas), dyes, and tanning-materials, would scarcely be in place in this volume.

The course through which the reader has been conducted will, it is hoped, have taught the principles of the science of chemistry, and given special knowledge concerning the applications of that science to medicine and pharmacy, as well as have imparted sufficient manipulative skill to meet the requirements of manufacture or analysis. The author would venture to suggest that this knowledge be utilized, not only in the way of personal advantage, but in experimental researches on chemical subjects connected with therapeutics

and pharmacy. The discovery and publication of a new truth, great or small, is the best means whereby to aid in advancing the calling in which we may be engaged, benefit our fellow creatures and ourselves, and contribute to that "ultimate end of knowledge" which Bacon defined as "the glory of the Creator and the endowment of human life."

INSTRUCTIONS TO BINDER.

PLATES.

"STARCHES" should face page 476.

"URINARY DEPOSITS, 1" should face page 597.

"URINARY DEPOSITS, 2" should face page 599.

APPENDIX.

TABLE OF OFFICIAL TESTS FOR IMPURITIES IN PREPARATIONS OF THE BRITISH PHARMACOPŒIA.*

Name of Preparation.	Impurities.	Test.	Page.
Acacise Gummi	Starch	Iodine	474
Acetum	More than one thousandth of Sulphuric acid	Quantitative analysis	701
Acidum Aceticum	Traces of Lead or Copper.	Sulphuretted Hydrogen	296
	Sulphuric acid	Chloride or Nitrate of Barium	362
Acid. Acetic. Glaciata	Hydrochloric acid	Nitrate of Silver	310
	Sulphurous acid	Nascent Hydrogen	357
Acidum Boracicum	Mineral matter	Evaporation and ignition	108
	Sulphurous acid	Nascent Hydrogen	357
Acidum Citricum	Alkaline salts	Insolubility in alcohol	390
	Traces of Copper or Lead.	Sulphuretted Hydrogen	296
Acidum Gallicum	Tartaric acid	Acetate of Potassium	376
	Sulphuric acid	Excess of Lime-water	381
Acidum Hydrochloricum	Mineral matter	Chloride or Nitrate of Barium	362
	Tannic acid	Incineration	108
Acidum Hydrocyanicum Dilutum	Sulphuric acid	(Isinglass) Gelatine	418
	Arsenic	Chloride or Nitrate of Barium	362
Acidum Nitricum	Sulphurous acid	Sulphuretted Hydrogen, Copper	196
	Sulphuric acid	Nascent Hydrogen	191
Acidum Oxallicum	Hydrochloric acid	Chloride or Nitrate of Barium	357
	Mineral matter	Ppt. by Nitr. silver, insol. in nitric acid	310
Acidum Phosphoricum Dilutum	Sulphuric acid	Evaporation and ignition	108
	Lead or Platinum	Chloride or Nitrate of Barium	362
Acidum Sulphuricum	Sulphuric acid	Nitrate of Silver	310
	Hydrochloric acid	Incineration	108
Acidum Tartaricum	Metaphosphoric acid	Sulphuretted Hydrogen	296
	Nitric acid	Chloride or Nitrate of Barium	362
Acidum Valerianicum	Phosphorous acid	Nitrate of Silver and Nitric acid	310
	Ammonium (phosphate)	Albumen	409
Acidum Valerianicum	Phosphorous acid	Ferrous Sulphate and Sulphuric acid	335
	Ammonium (phosphate)	Corrosive sublimate	411
Acidum Valerianicum	Phosphorous acid	Hot alkali	104
	Ammonium (phosphate)		

* The manipulations necessary to be observed in testing for impurities will be found described in the paragraphs treating of those substances. The Table also includes references to processes for ascertaining deficiency in strength of official articles.

The other "characters and tests" of pharmacopœial compounds have been given in connection with the respective synthetical and analytical reactions.

Name of Preparation.	Impurities.	Test.	Page.
Acidum Sulphuricum	Mineral matter	Evaporation and ignition	109
	Nitric acid	Ferrous Sulphate	335
	Arsenic or Lead	Sulphuretted Hydrogen	296
	Sulphuric acid	Chloride or Nitrate of Barium	362
Acidum Tannicum	Mineral matter	Incineration	109
	Metallic matter, as Lead	Sulphuretted Hydrogen	296
Acidum Tartaricum	Oxalic acid	Sulphate of Calcium	349
	Calcium (tartrate or sulphate).	Oxalate of Ammonium	124
Aconitia	Mineral matter	Incineration	109
	Mineral matter	Incineration	109
Adeps præparatus	Chloride (of Sodium)	Nitrate of Silver	310
	Starch (flour)	Iodine	474
Æther	Lime	Incineration, etc., as for gum	123
	Alcohol	Boiling-point, sp. gravity	511
Æther Aceticus	Water, Alcohol	Spec. grav. and boiling-point	513
Æther Purus	Alcohol, Water	Specific gravity	511
Alcohol	Resin or oil	Opalescence on dilution	506
	Water	Anhydrous sulphate of Copper	506
Alcohol Amylicum	Other spirituous matter	Boiling-point, sp. gravity	525
Alum	Iron (sulphate)	Yellow or red Prussiate	177
Ammonia Benzoas	Fixed salts	Non-volatility	106
	Fixed salts	Non-volatility	106
Ammonia Carbonas	Sulphate (of Ammonium)	Chloride or Nitrate of Barium	362
	Chloride (of Ammonium)	Nitrate of Silver	310
Ammonia Nitras	Chloride (of Ammonium)	Nitrate of Silver	310
	Sulphate (of Ammonium)	Chlor. or Nitrate of Barium	362
Ammonii Chloridum	Fixed salts	Non-volatility	106
Amylum	Alkaline matter	Red Litmus	101
	Acid matter	Blue Litmus	101
Antimonium Nigrum	Silica	Insol. in Hydrochloric acid	414
Antimonii Oxidum	Higher oxides of Antimony	Insolubility in sol. of acid tartrate of potassium	202
Antimonium Tartaratum	General	Quantitative analysis	698
Aqua Aurantii Floris	Metallic matter (Pb, Cu, Sn)	Sulphuretted Hydrogen	296
	Fixed salts	Evaporation and ignition	109
	Tin, Lead, or Copper	Sulphuretted Hydrogen	296
	Calcium salts	Oxalate of Ammonium	124
Aqua Destillata	Chlorides	Nitrate of Silver	310
	Sulphates	Chloride or Nitrate of Barium	362
Argenti Nitras	Carbonates	Lime-water	366
	Other nitrates, etc.	Quantitative analysis, etc.	694
Argenti Oxidum	Metallic silver	Effervescence with nitric acid	247
	General	Quantitative analysis	694
Argentum Purificatum	Copper	Ammonia to the Nitric solution	217
Asafetida	Earthy matter	Incineration	109
Atropa	Mineral matter	Incineration	109
Balsamum Peruvianum	Mineral matter	Incineration	109
	Fixed oil	Immiscibility with alcohol when mixed with water	109
Beberis Sulphas	Alcohol	Non-diminution of volume when mixed with water	109
	Mineral matter	Incineration	109
Bismuthi Carbonas	Nitrates (of Bismuth or Ammonium)	Sulphate of Indigo	337
	Lead (carbonate)	Diluted Sulphuric acid	244
	Chlorides (oxychloride of bismuth).	Nitrate of Silver	310

Name of Preparation.	Impurities.	Test.	Page.
Bismuthi oxidum	Moisture	Heated it loses weight.	708
	Carbonate (of Bismuth)	Sulphuric acid.	244
	Lead compounds	Nitrate of Silver	310
	Chlorides	HCl after sep. of Bi by AmHO	249
Bismuthi Subnitras	Silver	Diluted Sulphuric acid.	244
	Lead (oxynitrate).	Nitrate of Silver	310
	Chlorides (oxychloride of bismuth).		
Bismuthum Purificatum.	Copper.	Ammonia to Nitric solution	217
Borax	General	Quantitative analysis	649
Bromum	General	Specific gravity (830) and boiling-point	615
	Iodine	Starch.	317
Cadmii Iodidum	Zinc (iodide)	Potash in excess, then Sulphhydrate of Ammonium	298
	General	Quantitative analysis	698
Calcii Chloridum.	Hypochlorite of Calcium	Hydrochloric acid	340
Calcis Carbonas Precipitata	Carbonic acid	Lime-water	367
	Alumina, Oxide of Iron, and Phosphates.	Sacc. solution of lime to sol. in nitric acid	110
	Chlorides	Nitrate of Silver	310
Calcis Phosphas	Carbonates (of Calcium)	Effervescence with acids	308
	Alumina	Solution of potash, etc.	162
	Sand	Insolubility in acid	414
Calx	Carbonate of Calcium	Effervescence with acids	368
	Alumina, Oxide of Iron, etc.	Sacc. sol. of lime to sol. in acids	110
Calx Chlorata	General	Quantitative analysis	671
Cambogia	Starch	Iodine (Green).	474
Cambogia	Rice, flour, sand, powdered bark.	Microscope, after exhausting with spt. and with cold water	475
	Fixed salts	Non-volatility	106
Camphora	Earthy salts	Incineration (by help of red ox. of mercury)	108
Carbo Animalis Purificatus		Incineration	108
Carbo Ligni	More than 2 per cent. of ash	Iodine	474
Catechu Pallidum	Starch	Melting-point	616
Cera Alba	Soft Fats	Melting-point	616
	Soft Fats	Solubility in alcohol	560
Cera Flava	Resin	Insolubility in Turpentine, Iodine	474
	Flour	Ash sol. in acids with effervescence	368
Cerii Oxales	Carbonates and other oxalates.	Insolubility of hydrate in Ammonia	261
	Alumina	More or less than 46 per cent. of ash	261
	General	Melting-point	616
Cetaceum	Soft Fats	Litmus-paper	522
Chloral hydras	Free acid	Sulphuric acid.	524
	Other organic bodies	Quantitative analysis	523
	General	Specific gravity	630
Chloroform	General	Sulphuric acid.	521
	Hydrocarbons	Residue on evaporation	108
	Non-volatile matter	Lieben's test	506
Copaiba	Alcohol	Gelatinization at 270° F.	583
	Wood-oil	Incomplete solubility in benzol. (Not trustworthy.)	
Creasotum		Oxidation	529
	Carbolic acid	Non-volatility at 213° F.	529
		Dextro-rotation of polarized ray	529
		Crystallization on cooling	529

Name of Preparation.	Impurities.	Test.	Page.
Cupri Sulphas . . .	Iron (ferrous sulphate) . .	Nitric acid and Ammonia . .	177
Elaterium . . .	Carbonates (chalk) . . .	Effervescence with acids . .	366
Fel Bovinum Purificatum . . .	General . . .	Quantitative analysis . . .	466
	Mucous, crude bile . . .	Incomplete solubility in spirit . . .	546
Ferri Arsenias . . .	Sulphate (of Sodium) . . .	Chloride or Nitrate of Barium . .	363
	General . . .	Quantitative analysis . . .	667
Ferri Carbonas Saccharata . . .	Sulphate (of Ammonium) . .	Chloride or Nitrate of Barium . .	368
	General . . .	Quantitative analysis . . .	668
	Tartrate (of Iron and Ammonium) . .	Ebullition with potash and saturation with acetic acid (= $\text{KHC}_2\text{H}_3\text{O}_6$) . .	376
Ferri et Ammonis Citras . . .	General . . .	Quantitative analysis . . .	696
	Potassium or Sodium (salts) . .	Alkalinity of ash . . .	101
	Potassium or Sodium (salts) . .	Alkalinity of ash . . .	101
Ferri et Quinise Citras . . .	General . . .	Quantitative analysis . . .	732
	Alkaloids other than Quinia . .	Insolubility of pptd. alkaloid in ether . . .	456
Ferri Oxidum Magneticum . . .	Metallic iron . . .	Effervescence with acids . .	173
	General . . .	Quantitative analysis . . .	686
Ferri Peroxidum Humidum . . .	Ferrous hydrate . . .	Red prussiate to acid solution . . .	177
	Ferric oxyhydrate . . .	Insolubility in cold dil. Hydrochloric acid . .	166
	Arsenicum (ferri arsenias) . .	Slip of Copper in acid solution . . .	191
Ferri Phosphas . . .	General . . .	Quantitative analysis . . .	667
	Ferric oxysulphate . . .	Insolubility in water . . .	167
Ferri Sulphas Granulata . . .	Ferric compounds . . .	Ppt. of Sulphur in aq. sol. by sulphuretted hydrogen . .	176
	Copper, etc.	Sulphuretted Hydrogen . . .	296
Ferrum Redactum . . .	Less than 50 per cent. . .	Quantitative analysis . . .	667
	Ferrous compounds . . .	Red prussiate to acid solution . . .	177
Ferrum Tartaratum . . .	Ammoniacal salts . . .	Soda . . .	101
	General . . .	Quantitative analysis . . .	686
	General . . .	Specific gravity . . .	630
Glycerinum . . .	Fixed Salts . . .	Non-volatility . . .	236
Hydrargyri Iodidum Rubrum . . .	Red Iodide . . .	Solubility in ether . . .	221
Hydrargyri Iodidum Viride . . .	Fixed salts, as NaCl . . .	Non-volatility . . .	236
Hydrargyri Oxidum Flavum . . .	Fixed salts . . .	Non-volatility . . .	236
Hydrargyri Oxidum Rubrum . . .	Nitrates (of Mercury) . .	Orange vapour on heating in a tube . . .	230
Hydrargyri Subchloridum . . .	Corrosive Sublimato . . .	Treatment with ether . . .	239
Hydrargyri Sulphas . . .	Fixed salts . . .	Non-volatility . . .	236
Hydrargyrum . . .	Fixed salts . . .	Non-volatility . . .	236
	Fixed metals (Pb, Sn, Zn, Bi, Cu) . . .	Non-volatility . . .	236
Hydrargyrum Ammoniatum . . .	Fixed salts . . .	Non-volatility . . .	236
Hydrargyrum cum Creta . . .	Mercuric oxide . . .	Stannous chloride to solution in hydrochloric acid . .	236
	Fixed salts . . .	Non-volatility . . .	315
Iodum	Cyanide of Iodine . . .	Physical characters . . .	315
	General . . .	Quantitative analysis . . .	671
Jalape Resina . . .	Resin . . .	Solubility in turpentine . . .	467
Limonis Succus . . .	Deficiency of Citric acid . .	Quantitative analysis . . .	379
Liquor Ammonis . . .	General . . .	Specific gravity . . .	630
		Quantitative analysis . . .	647

Name of Preparation.	Impurities.	Test.	Page.
Liquor Ammoniae Fortior.	General impurity or deficiency.	Specific gravity (630) and Quantitative analysis . . .	647
	Carbonate of Ammonium.	Lime-water	366
	Calcium salts (chloride etc.)	Oxalate of Ammonia	124
	Iron salts (ferrous hydrate).	Sulphydrate of Ammonium .	177
	Sulphur salts (AmHS) . .	Ammonio-sulphate of Copper	364
Liq. Antimonii Chloridi	Chloride of Ammonium . .	Nitrate of Silver to acidified sol.	310
	Sulphate of Ammonium . .	Chloride of Barium to acidified sol.	362
			630
	General impurity or deficiency	Specific gravity	630
		Quantitative analysis	664
Liq. Arsenicæ			
Liq. Arsenici Hydrochloricus . .			
Liq. Bismuthi et Ammoniae Citæ			690
Liq. Calcis	Deficiency in strength . .	Quantitative analysis	660
Liq. Calcis Chloratæ	General impurity or deficiency	Specific gravity	630
Liq. Calcis Saccharatus		Quantitative analysis	660
Liquor Chlori	General quality	Specific gravity	630
	Fixed matter	Residue on evaporation . . .	108
	Deficiency in strength . .	Quantitative analysis	671
Liq. Ferri Perchloridi Fort. . .			
Liq. Ferri Pernitratæ	Ferrous salts	Red prussiate	177
Liq. Ferri Persulphatis	General impurity or deficiency	Specific gravity	630
		Quantitative analysis	666
Liq. Hydrargyri Nit. Acid	Deficiency in strength . .	Specific gravity	630
Liq. Lithiæ Efferves.	Mercurous salts (nitrate) .	Hydrochloric acid	235
Liq. Magnesiæ Carbonatæ . .	Gen. impur. or deficiency .	Quantitative analysis, etc. . .	269
Liq. Plumbi Subacetatis	Other magnesian salts . .	Bitter taste (MgCl, MgSO ₄) .	
	Gen. impur. or deficiency .	Quantitative analysis	633
	General impurity or deficiency	Specific gravity	630
		Quantitative analysis	660
	General impurity or deficiency	Specific gravity	630
		Quantitative analysis	664
	Carbonate of (Potassium) .	Effervescence with acids . . .	366
Liquor Potassæ	Calcium salts	Lime-water	366
		Oxalate of Ammonium	124
	More than traces of { Silica	Insol. in acid after evap. etc.	414
	{ Sulphates	Chloride of Barium to acid sol.	362
		Nitrate of Silver to acid sol..	310
Liq. Potassæ Effervescens	Chlorides	Ammonia to acid sol.	153
	Alumina	Quantitative analysis	661
	Deficiency in strength . .	Tartaric acid, etc.	678
	Bicarbonate of Sodium . .	Specific gravity	630
	General impurity or deficiency	Quantitative analysis	661
Liq. Sodæ	Calcium salts	Oxalate of Ammonium	124
	Carbonate (of sodium) . .	Effervescence with acids . . .	366
		Lime-water	366
	More than traces of { Silica	Insol. in acids after evap. etc.	414
	{ Sulphates	Barium chloride to acid sol.	362
Silver nitrate to acid sol. . .		310	
	Alumina	Ammonia to acid sol.	153
Liquor Sodæ Chloratæ	Salts of Potassium or Ammonium.	Perchloride of Platinum to acid sol.	79, 101
	General impurity or deficiency.	Quantitative analysis	661
	Calcium salts	Oxalate of Ammonium	124

Name of Preparation.	Impurities.	Test.	Page.
Liq. Sodæ Effervescent	Deficiency in strength . . .	Quantitative analysis . . .	651
	General impurity or deficiency.	Quantitative analysis . . .	259
Lithiæ Carbonas . . .	Calcium salts	Oxalate of Ammonium, etc. . .	259
	Alumina	Lime-water, etc.	259
Lithiæ Citras	Deficiency in strength . . .	Quantitative analysis . . .	259
	Starch, etc.	Microscope	475
Lycopodium	More than 5 per cent. of mineral matter . . .	Incineration	108
	Carbonate (of Magnesium)	Earthy matter sinks in CS ₂ . .	
Magnesia	Calcium (hydrate or carbonate).	Effervescence with acids . . .	129
Magnesia Levis	Sulphates (of Magnesium or Sodium).	Oxalate of Ammonium to acet. sol.	124
	Alumina	Chloride of Barium to acid solution	362
Magnesii Carbonas . .	Sulphates (of Magnesium or Sodium).	Ammonia to acid sol.	152
Magnesii Carbonas Levis	Calcium (carbonate).	Chloride of Barium to acid solution	362
	Iron, Lead, etc.	Oxalic Acid to ammoniacal solution	134
	General impurity or deficiency	Sulph. Hydrogen to sol. in acid + excess of Ammon. . .	296
	Calcium (sulphate) . . .	Quantitative analysis . . .	663
Magnesii Sulphas . . .	Iron (sulphate)	Oxalate of Ammonium . . .	124
	General impurities . . .	Chlorinated Lime or Soda . .	127
Manna	Deficiency of mannite . . .	Quantitative analysis . . .	683
Mel	Starch (flour)	Quantitative analysis . . .	444
Morphiæ Hydrochloras . .	General impurities	Iodine	474
		Quantitative analysis . . .	723
Opium	Deficiency in Morphia . . .	Quantitative analysis . . .	723
Plumbi Acetas	General	Quantitative analysis . . .	682
	Sulphate of Lead or Barium or Silicates.	Insolubility in Acetic acid . .	
Plumbi Carbonas	Calcium (chalk)	110, 244, Oxal. Ammonium after removing Lead	414
	Carbonates	Effervescence with acids . . .	252
Plumbi Oxidum	Copper (oxide)	Ammonia to acid solution . .	217
	More than { Chloride . . .	Nitrate of Silver to acid sol. .	310
	traces of { Sulphate . . .	Chloride of Barium to acid sol.	362
Potassæ Caustica	General impurities (water etc).	Quantitative analysis . . .	651
Potassæ Sulphurata . . .	Excess of Carbonate or Sulphate	More than 75 per cent. in sol. in spirit	2
Potassæ Acetas	Iron and other metallic impurities	Sulphydrate of Ammonium . .	29
	Carbonate of Potassium . .	Effervescence with acids; alkalinity; insolubility in spirit.	396
Potassæ Bicarbonas . . .	General	Quantitative analysis . . .	651
	More than { Silicate . . .	Insol. in acids after evap. etc.	414
Potassæ Carbonas	traces of { Sulphate . . .	Chloride of Barium to acid sol.	362
	{ Chloride	Nitrate of Silver to acid sol. .	310
	General	Quantitative analysis . . .	651
Potassæ Chloras	Chloride (of Potassium)	Nitrate of Silver	310
	Calcium (chloride)	Oxalate of Ammonium . . .	124
Potassæ Citras	General	Quantitative analysis . . .	651
Potassæ Nitras	Sulphate (of Potassium)	Chloride of Barium	362
Potassæ Permanganas . .	Chloride (of Potassium)	Nitrate of Silver	310
	General	Quantitative analysis . . .	651

Name of Preparation.	Impurities.	Test.	Page.
Potassæ Sulphas . . .	Acid Sulphate of Potassium	Test-paper	101
Potassæ Tartaras . . .	Calcium (sulphate) . . .	Oxalate of Ammonium . . .	124
Potassæ Tart. Acida . . .	General	Quantitative analysis . . .	653
Potassii Bromidum . . .	Free Bromine	Odour	311
Potassii Ferridcyanidum . . .	Iodide (of Potassium) . . .	Chlorine-water and starch . . .	317
	General	Quantitative analysis . . .	661, 696
	Ferrocyanide (of Potassium).	Ferric salt	398
Potassii Iodidum . . .	Iodate of (Potassium) . . .	Tartaric acid and starch . . .	77
	Chloride (of Potassium) . . .	Nitrate of Silver, etc. . . .	317
	Carbonate (of Potassium) . . .	Sacc. solution of Lime . . .	386
Quinæ Sulphas . . .	Salicin	Sulphuric acid	493
Rhei Radix . . .	General	Quantitative analysis . . .	720
Santoninum . . .	Turmeric	Boracic acid	591
	Mineral matter	Incineration	108
Sapo Durus . . .	Earthy soaps, etc.	Insolubility in spirit	544
	Oil	Oily stain to paper	545
	Potassium compounds . . .	Deliquescence of ash	545
Sapo Mollis . . .	Earthy soaps, etc.	Insolubility in spirit	544
	Oil	Oily stain to paper	545
	Resin of guaiacum	Inner surface of Potato-paring	497
Scammonis Resina . . .	Resin of Jalap	Insolubility in ether	500
	Carbonate (of Calcium and Magnesium).	Effervescence with acids . . .	366
Scammonium . . .	Starch (flour)	Solution of Iodine	474
	General impurities	Quantitative analysis	500
Sinapis . . .	Starch (flour)	Solution of Iodine	474
	More than { Chloride	Nitrate of Silver to acid sol. . .	310
	traces of { Sulphate	Chloride of Barium to acid sol. . .	362
Soda Caustica . . .	General impurities (water etc.)	Quantitative analysis	651
Soda Tartarata . . .	General	Quantitative analysis	652
	Acid or alkaline matter . . .	Test-paper	101
Sodæ Acetas . . .	Sulphates (Sodium or Calcium).	Chloride of Barium to acid sol. . .	362
	Chlorides (Sodium or Calcium).	Nitrate of Silver to acid sol. . .	310
Sodæ Arsenias . . .	Excess or deficiency of water of crystallisation.	Quantitative analysis	708
	General	Quantitative analysis	661
	Neutral Carbonate (of Sodium).	Mercuric Chloride	236
Sodæ Bicarbonas . . .	More than { Chlorides	Nitrate of Silver to acid sol. . .	310
	traces of { Sulphates	Chloride of Barium to acid sol. . .	362
	General	Quantitative analysis	651
	Excess or deficiency of water of cryst., etc. . .	Quantitative analysis	608
Sodæ Carbonas . . .	More than { Chlorides	Nitrate of Silver to acid sol. . .	310
	traces of { Sulphates	Chloride of Barium to acid sol. . .	362
Sodæ Hyposulphis . . .	General	Quantitative analysis	665
Sodæ Nitras . . .	Chloride (of Sodium)	Nitrate of Silver	310
	Sulphate (of Sodium)	Chloride or Nitrate of Barium	362
Sodæ Phosphas . . .	More than trace of Sulphate	Chloride or Nitrate of Barium to acid sol.	362
	Deficiency or excess of water of cryst.	Quantitative analysis	708

Name of Preparation.	Impurities.	Test.	Page.
Sodæ Sulphas . .	Ammonium salts	Solution of potash heated. { Quantitative analysis Quantitative analysis	104
	Ferric salts		178
	General		701
	Excess or deficiency of water of cryst.		708
Sodæ Valerianas . .	Free soda or carbonate . .	Test-papers Insolubility in spirit Specific gravity	101
	General		424
	More than trace of acid . .		630
Spiritus Ætheris Nitrosi	Free acid	Effervescence with bicarbonate of sodium More than "feeble" effervescence with bicarbonate of sodium	368
			368
			513
Spiritus Ammoniac-Aromaticus . .	General	Quantitative analysis	513
			513
Spiritus Chloroformi	General	Specific gravity	630
Spiritus Rectificatus	General (excess of water) .	Specific gravity	630
	Resin or oil	Opalescence on dilution . . .	546
	More than traces of fusel oil, etc.	Nitrate of Silver	662
Spiritus Tenuior	General (excess of water) .	Specific gravity	508
Strychnia	Brucia	Nitric acid	461
	Mineral matter	Incineration	106
Sulphur Præcipitatum	Sulphate of Calcium	Appearance under microscope Residue on ignition Incineration	363
			354
			106
Sulphur Sublimatum	Earthy matter	Litmus-paper	101
	Trace of acid (H_2SO_4 or H_2SO_3) .		
Sulphuris Iodidum Symplic	Sulphide of Arsenicum	Ammonia	198
	Deficiency of Iodine	Quantitative analysis	316
Tamarindus	Deficiency of Sugar	Specific gravity	630
	Traces of Copper	Iron	216
Veratria	Mineral matter	Incineration	106
	Sulphates	Chloride or Nitrate of Barium	362
Zinci Acetas	Chlorides	Nitrate of Silver	310
	Metals (As, Cd, Cu, Pb) . .	Sulphuretted Hydrogen	296
	Iron (acetate)	Nitric acid; Ammonia	140
	Copper (acetate)	Ammonia	217
	Sulphates	Chloride or Nitrate of Barium to acid sol.	362
Zinci Carbonas	Chlorides	Nitrate of Silver to acid sol. .	310
	Copper (carbonate)	Ammonia to acid solution . .	217
	Metals (As, Cd, Cu, Pb) . .	Sulphuretted Hydrogen	296
	Sulphates	Chloride or Nitrate of Barium	362
Zinci Chloridum	Calcium (chloride)	Oxalate of Ammonium	124
	Ferrous salts (chloride) . . .	Ferridcyanide of Potassium . .	177
	Ferric salts (chloride)	Ferrocyanide of Potassium . .	174
	Carbonate (of Zinc)	Effervescence with acids . . .	368
	Sulphates (Sodium or Zinc) .	Chloride or Nitrate of Barium to acid sol.	362
Zinci Oxidum	Chlorides	Nitrate of Silver to acid sol. .	310
	Copper (oxide)	Ammonia to acid solution . .	217
	Metals (As, Cd, Cu, Pb) . .	Sulphuretted Hydrogen	310
	Iron (sulphate)	Tincture of galls	414
Zinci Sulphas	Copper (sulphate)	Nitric acid; Ammonia	140
		Ammonia	217
Zinci Valerianas	Sulphate (of Zinc)	Chloride or Nitrate of Barium	362
	Butyrate (of Zinc)	Acetate of Copper, etc.	424

SATURATION TABLES.

Equivalent weights of Citric Acid, Tartaric Acid, Carbonate of Potassium, Bicarbonate of Potassium, Carbonate of Sodium, Bicarbonate of Sodium, Carbonate of Ammonium, and Carbonate of Magnesium; repeated (in black) for 20 parts of each, and incidentally (in roman) for other proportions. (Exact to two places of decimals.)

Citric Acid ($H_3C_6H_5O_7$, H_2O) $\div 3 \times 2 = 140$	20.00	18.66	16.96	14.00	9.78	16.66	23.72	29.31
Tartaric Acid $H_2C_4H_4O_6 = 150$	21.43	20.00	18.26	15.00	10.49	17.85	25.42	31.41
Carbonate of Potassium $K_2CO_3 + 16\% Aq = 164.285$	23.47	21.90	20.00	16.43	11.48	19.52	27.87	34.40
Bicarbonate of Potassium $2(KHCO_3) = 200$	28.57	26.66	24.34	20.00	13.98	23.81	33.99	41.90
Carbonate of Sodium Na_2CO_3 , $10H_2O = 286$	40.08	38.13	34.81	28.60	20.00	34.04	48.47	59.98
Bicarbonate of Sodium $2(NaHCO_3) = 168$	24.00	22.40	20.45	16.80	11.74	20.00	28.47	35.18
Carbonate of Ammonium $(N_4H_{16}C_3O_6) \div 2 = 118$	16.85	15.73	14.36	11.80	8.25	14.04	20.00	24.71
Carb. of Magnes. $(MgCO_3)_2, Mg_2HO, 4H_2O \div 4 = 96.5$	13.64	12.73	11.62	9.55	6.68	11.37	16.18	20.00

The amount of acid given in any column will saturate the amount of carbonate in the same column, and *vice versa*.

The amounts of carbonate in any column are equal to each other in chemical power.
Lemon-Juice (sp. gr. 1.039) contains, on an average, 7 per cent. by weight of citric acid.

The same Table in round numbers, for purposes of prescribing and dispensing.
(The old names in Latin.)

Citric Acid	20	19	17	14	10	17	24	30
Tartaric Acid	22	20	18	15	11	18	26	32
Carbonate of Potassium (Potassæ Carbonas, B. P.)	24	22	20	16	12	20	28	35
Bicarbonate of Potassium (Potassæ Bicarbonas, B. P.)	28	27	24	20	14	24	34	42
Carbonate of Sodium (cryst.) (Sodæ Carbonas, B. P.)	40	38	34	28	20	34	49	60
Bicarbonate of Sodium (Sodæ Bicarbonas, B. P.)	24	23	20	17	13	20	29	36
Carbonate of Ammonium (Ammonis Carbonas, B. P.)	17	16	14	12	8	14	20	25
Carbonate of Magnesium (Magnesiæ Carbonas, B. P.)	13	13	11	9	7	11	16	20

The Table is read thus:—30 grains of Citric Acid will saturate 29 grains of Bicarbonate of Potassium; 20 grains of Bicarbonate of Sodium will saturate, &c. &c. 15 grains of Tartaric Acid; 11 grains of Tartaric Acid = 8 grains of Carbonate of Ammonium; 20 grains of Bicarbonate of Sodium are equivalent to, or will do as much work as, 34 grains of Carbonate of Sodium; 14 grains of Citric Acid are as strong as 15 of Tartaric Acid. It is occasionally convenient to double the numbers, halve them, or take some other proportion; also to employ them in weights other than grains.

Lemon-Juice contains, on an average, 32½ grains of Citric Acid in 1 fluid ounce, or 4 grains per fluid drachm.

THE ELEMENTS.

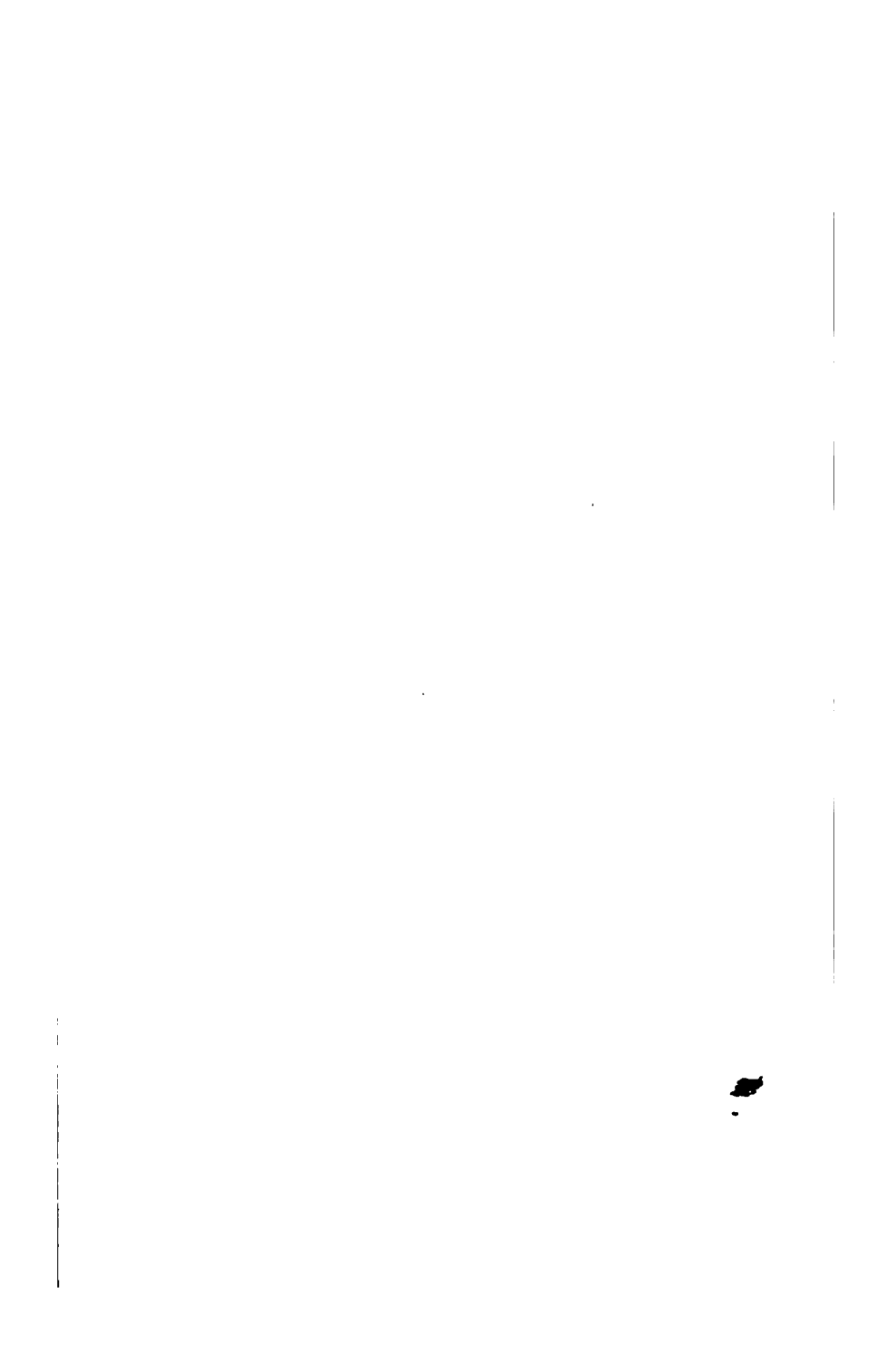
	Symbol and atomic value.	Atomic weight.
Aluminium (Al^{VI})	Al^{IV}	27.5
Antimony (Sb^{III})	Sb^{V}	122
Arsenicum (As^{III})	As^{V}	75
Barium	Ba^{II}	137
Beryllium (or Glucinum)	Be^{II}	9.5
Bismuth (Bi^{III})	Bi^{V}	208
Boron	B^{III}	11
Bromine ($^{79.75}$, Stas)	Br^{I}	80
Cadmium	Cd^{II}	112
Cæsium	Cs^{I}	133
Calcium	Ca^{II}	40
Carbon (C^{II})	C^{IV}	12
Cerium (Ce^{II}) (138 , Mendelejeff)	Ce^{VI}	92
Chlorine ($^{35.45}$, Stas)	Cl^{I}	35.5
Chromium (Cr_2^{VI})	Cr^{VI}	52.5
Cobalt (Co^{II})	Co^{VI}	58.8
Copper	Cu^{II}	63.5
Davyum	...	...
Decipium	...	...
Didymium (138 , Mendelejeff)	D^{II}	96
Erbium (P) (171 , Mendelejeff)	Er^{II}	112.6
Fluorine	F^{I}	19
Gallium	...	...
Glucinum, <i>see</i> Beryllium	...	...
Gold (Au)	Au^{III}	196.7
Hydrogen	H^{I}	1
Indium (113 , Mendelejeff)	In^{VI}	75.6
Iodine ($^{126.93}$, Stas)	I^{I}	127
Iridium	Ir^{IV}	197
Iron (Fe^{II} & Fe_2^{VI})	Fe^{VI}	56
Lanthanium (138 , Mendelejeff)	La^{II}	92
Lavæsium	...	...
Lead (Pb^{II})	Pb^{IV}	207
Lithium ($^{7.004}$, Stas)	Li^{I}	7
Magnesium	Mg^{II}	24
Manganese (Mn^{II} & Mn^{IV})	Mn^{VI}	55
Mercury	Hg^{II}	200
Molybdenum	Mo^{VI}	96
Mosandrum	...	...
Neptunium	...	...
Nickel (Ni^{II})	Ni^{VI}	58.8

THE ELEMENTS (*continued*).

	Symbol and atomic value.	Atomic weight.
Niobium	Nb ^V	97.6
Nitrogen (N ^I & N ^{III}) (14.008 Stas)	N ^V	14
Norwegium	...	...
Osmium	Os ^{IV}	199
Oxygen (15.96 Stas)	O ^{II}	16
Palladium	Pd ^{IV}	106.5
Phillipium	...	...
Phosphorus (P ^{III})	P ^V	31
Platinum (197.23, Andrews)	Pt ^{IV}	198
Potassium (39.04 Stas)	K ^I	39
Rhodium	Rh ^{IV}	104.3
Rubidium	Rb ^I	85.3
Ruthenium	Ru ^{IV}	104.2
Scandium	...	...
Selenium (or Selenion)	Se ^{VI}	79.5
Silicon	Si ^{IV}	28
Silver (107.86 Stas)	Ag ^I	108
Sodium (22.96 Stas)	Na ^I	23
Strontium	Sr ^{II}	87.5
Sulphur (S ^{II} & S ^{IV})	S ^{VI}	32
Tantalum	Ta ^V	182
Tellurium	Te ^{VI}	129
Terbium (P)	...	...
Thallium (208, Crookes)	Tl ^{III}	204
Thorium (or Thorium)	Th ^{II}	232
Tin (Sn ^{II})	Sn ^{IV}	118
Titanium	Ti ^{IV}	50
Tungsten	W ^{VI}	184
Uralium (187.25, Guyard)	...	...
Uranium (240, Mendelejeff)	U ^{VI}	120
Vanadium	V ^V	51.3
Ytterbium	...	...
Yttrium (90, Mendelejeff)	Y ^{III}	61.7
Zinc	Zn ^{II}	65
Zirconium	Zr ^{IV}	89.5

The quantivalence or atomic value of some elements is, apparently, variable: in the above Table the full coefficients are given in the column of symbols, other common values in brackets.

Atomic weights are sometimes obscurely termed equivalents.



INDEX.

- Abies balsamea*, 557, 564.
 „ *excelsa*, 562.
Abrus precatorius, 496.
Absinthol, 554.
Absolute alcohol, 501, 629.
Acacia catechu, 419.
 „ *suma*, 419.
Acacia gummi, 123.
 „ *gummi*, impurities in, 735.
Acetate of ammonium, solution of, 97.
 „ *amyl*, 525.
 „ *copper*, 216.
 „ *ethyl*, 349, 513.
 „ *iron*, 165, 349.
 „ *lead*, 239.
 „ *morphia*, 449.
 „ *potassium*, 69.
 „ *sodium*, 85.
 „ *zinc*, 145.
Acetates, 345.
 „ *analytical reactions of*, 349.
 „ *decomposition of aqueous solutions of*, 347.
 „ *volumetric estimation of*, 619.
Acetic acid, 345.
 „ *glacial*, 347, 592.
 „ *volumetric estimation of free*, 619.
Acetic ether, 349.
Acetone, 349.
Acetonitrile of barium, 187.
 „ *iron*, 174.
Acetum, 346.
 „ *cantharidis*, 346.
 „ *impurities in*, 735.
 „ *scilla*, 346.
Acetyl, 346.
Acid, *acetic*, 345, 549, 629.
 „ *acetic, glacial*, 347, 629.
 „ *aconitic*, 379.
 „ *amyric*, 504.
 „ *angelic*, 553.
 „ *arabic*, 479.
 „ *arachidic*, 549.
Acid, *arsenic*, 187.
 „ *arsenious*, 186.
 „ *benzoic*, 392.
 „ *boracic*, 389.
 „ *bromic*, 78, 343.
 „ *butyric*, 424, 549.
 „ *camphoretic*, 558.
 „ *camphoric*, 558.
 „ *cantharidic*, 558.
 „ *caproic*, 549, 547.
 „ *caprylic*, 549, 547.
 „ *carbazotic*, 530.
 „ *carbolic*, 528.
 „ *carbonic*, 22, 364.
 „ *carminic*, 394.
 „ *catechuic*, 419.
 „ *cathartic*, 492.
 „ *cathartogenic*, 492.
 „ *cerotic*, 547, 549.
 „ *cetraric*, 394.
 „ *chavicic*, 470.
 „ *chloric*, 122, 339.
 „ *cholalic*, 545.
 „ *chromic*, 272.
 „ *chrysammic*, 530.
 „ *chrysophanic*, 394.
 „ *cinnamic*, 394, 566.
 „ *citric*, 378.
 „ *colophonic*, 561.
 „ *colophonic*, 561.
 „ *copaivic*, 563.
 „ *cresotic*, 531.
 „ *cresylic*, 529, 530.
 „ *cryptophanic*, 554.
 „ *cuminic*, 555.
 „ *cyanic*, 395.
 „ *dextroracemic*, 373.
 „ *dextrotartaric*, 373.
 „ *dithionic*, 406.
 „ *engenic*, 554.
 „ *equisetic*, 379.
 „ *erucic*, 549.
 „ *ethylsulphuric*, 505, 510.
 „ *ferulalic*, 564.
 „ *filicic*, 548.
 „ *fluoric*, 400.

- Acid, formic, 395, 549.
 „ gallic, 420.
 „ gambogic, 565.
 „ gaultheric, 526.
 „ gelseminic, 468.
 „ gentianic, 495.
 „ gentisic, 496.
 „ glacial acetic, 347.
 „ glacial phosphoric, 335.
 „ glycollic, 533.
 „ guaiaretinic, 496.
 „ gummic, 479.
 „ hemidesmic, 396.
 „ hippuric, 396, 598.
 „ hydriodic, 314.
 „ hydrobromic, 311.
 „ hydrochloric, 21, 305.
 „ hydrochloric, common, 305.
 „ hydrochloric, dilute, 305.
 „ hydrocyanic, 322, 491.
 „ hydrocyanic, dilute, 324.
 „ hydroferridecyanic, 399.
 „ hydroferrocyanic, 397.
 „ hydrofluoric, 400.
 „ hydrosulphuric, 350.
 „ hypochlorous, 122, 329.
 „ hypophosphoric, 412.
 „ hypophosphorous, 401.
 „ hyposulphurous, 404.
 „ iodic, 344.
 „ jalapic, 497.
 „ lactic, 406, 533.
 „ laevoracemic, 373.
 „ laevotartaric, 373.
 „ larixinic, 420.
 „ lauric, 549.
 „ lithic, 422.
 „ lupulinic, 564.
 „ malic, 407.
 „ mastichic, 562.
 „ meconic, 407.
 „ melissic, 549.
 „ metaboracic, 389.
 „ metagummic, 479.
 „ metantimonie, 203.
 „ metaphosphoric, 385, 408.
 „ metastannic, 278.
 „ mimotannic, 419.
 „ mucic, 489.
 „ myristic, 549.
 „ myrrhic, 565.
 „ naphthalic, 390.
 „ nitric, 329.
 „ nitric, dilute, 333.
 „ nitrohydrochloric, 209, 334.
 „ nitrohydrochloric, dilute, 209, 334.
 „ nitrous, 335, 401.
 Acid, osanthylic, 549.
 „ oleic, 544.
 „ ophelic, 410.
 „ opianic, 450.
 „ orthophosphoric, 410.
 „ oxalic, 368.
 „ oxalic, chemically pure, 369.
 „ oxybutyric, 533.
 „ palmitic, 542.
 „ paratartaric, 373.
 „ parietinic, 394.
 „ pelargonic, 549.
 „ pentathionic, 405.
 „ perchloric, 122, 342.
 „ phenic, 523.
 „ phosphomolybdic, 387, 538.
 „ phosphoric, 15, 383.
 „ phosphoric, dilute, 384.
 „ phosphorous, 385, 410.
 „ phthalic, 390.
 „ picric, 530.
 „ pimanic, 561.
 „ pinic, 560.
 „ polygalic, 500.
 „ propionic, 549.
 „ prussic, 322.
 „ pyrethric, 562.
 „ pyrogallic, 421.
 „ pyroligneous, 345.
 „ pyrophosphoric, 386, 412.
 „ racemic, 373.
 „ rheic, 395.
 „ rhubarbaric, 395.
 „ rutic, 549.
 „ saccharic, 489.
 „ salicylic, 526, 530.
 „ salicylous, 526.
 „ santonic, 499.
 „ sarcolactic, 407.
 „ sclerotic, 561.
 „ sclerotinic, 561.
 „ silicic, 413, 415.
 „ solution of arsenic, 185, 186.
 „ stannic, 279.
 „ stearic, 544, 549.
 „ succinic, 415.
 „ sulphethylic, 510.
 „ sulphindigotic, 337.
 „ sulphindyllic, 337.
 „ sulphocarbolic, 530.
 „ sulphocyanic, 416.
 „ sulphophenic, 530.
 „ sulphosalicylic, 531.
 „ sulphovinic, 361, 510.
 „ sulphuric, 358.
 „ sulphuric, aromatic, 361.
 „ sulphuric, dilute, 361.
 „ sulphurous, 21, 355.

- Acid, sulphydric, 350.
 „ sylvic, 560.
 „ tannic, 417.
 „ tartaric, 371.
 „ tetrathionic, 405.
 „ tiglic, 548.
 „ trinitrocarbohic, 530.
 „ trithionic, 405.
 „ tropic, 464.
 „ uric, 421, 595.
 „ valerianic, 423, 549.
 „ vanillic, 425.
 Acid carbonate of potassium, 71.
 „ carbonate of sodium, 85.
 „ salts, 73, 74, 852.
 „ solution of arsenic, 186.
 „ tartrate of potassium, 61, 80, 372.
 Acidimetry, 658.
 Acids, analytical detection of, 431.
 „ antidotes to, 305.
 „ definition of, 302.
 „ free, estimated, 655.
 „ of chlorine, 342.
 „ of phosphorus, 410.
 „ quantitative estimation of, 655.
 Acidulous radicals, formulæ and quantivalence of, 66, 133, 302, 303.
 „ radicals, qualitative detection of, 426.
 „ radicals, quantitative estimation of salts of, 695.
 „ radicals, Tables to aid in the detection of, 430, 431.
 Acidum aceticum, 847.
 „ aceticum, impurities in, 735.
 „ aceticum dilutum, 847.
 „ aceticum glaciale, 847.
 „ aceticum glaciale, impurities in, 735.
 „ arseniosum, 186.
 „ benzoicum, 392.
 „ boracicum, impurities in, 735.
 „ carbolicum, 528.
 „ chromicum, 272.
 „ citricum, 378.
 „ citricum, impurities in, 735.
 „ gallicum, 420.
 „ gallicum, impurities in, 735.
 „ hydriodicum dilutum, 315.
 „ hydrochloricum, 21, 305.
 „ hydrochloricum, impurities in, 735.
 „ hydrochloricum dilutum, 305.
 Acidum hydrocyanicum dilutum, 324.
 „ hydrocyanicum dilutum, impurities in, 735.
 „ nitricum, 333.
 „ nitricum, impurities in, 693.
 „ nitricum dilutum, 333.
 „ nitro-hydrochloricum dilutum, 209, 334.
 „ ozalicum, impurities in, 735.
 „ phosphoricum dilutum, 384.
 „ phosphoricum dilutum, impurities in, 735.
 „ sulphuricum, 361.
 „ sulphuricum, impurities in, 736.
 „ sulphuricum aromaticum, 361.
 „ sulphuricum dilutum, 361.
 „ sulphurosum, 356.
 „ sulphurosum, impurities in, 736.
 „ tannicum, 417.
 „ tannicum, impurities in, 736.
 „ tartaricum, 373.
 „ tartaricum, impurities in, 736.
 Acipenser, 540.
 Aconiti ferocis radix, 463.
 „ folia, 462.
 „ heterophylli radix, 463.
 „ radix, 462.
 Aconitia, 462.
 „ impurities in, 736.
 Aconitic acid, 379.
 Aconitina, 462.
 Aconitine, 462.
 Aconitum ferox, 463.
 „ heterophyllum, 463.
 „ napellus, 462.
 Acorin, 556.
 Acorus calamus, 556.
 Acridyl sulphocyanate, 532.
 Acrolein, 534.
 Actea racemosa, 473.
 Adeps benzoatus, 546.
 „ præparatus, 546.
 „ præparatus, impurities in, 736.
 Adhesion, 55.
 Adipocere, 540.
 Advice to students, xiii.
 Ægle Marmelos, 419.
 Aerated bread, 486.
 „ water, 88.
 Æsculin, 468.
 Ether, 509.
 „ impurities in, 726.
 „ aceticus, 513.
 „ aceticus, impurities in, 736.

- Æther, purus*, 511.
 „ *purus*, impurities in, 738.
 Affinity, chemical, 31.
 „ units of, 54, 132.
 Agate, 413.
 Air, composition of, 17.
 „ influence of animals and plants on, 9.
 „ nitrogen in the, 16.
 „ oxygen in the, 4.
 „ relative weight of the, 17.
 „ weight of 1 cubic cent., 636.
 „ weight of 100 cub. inches, 636.
 Ajowan oil, 552.
 Ajwain oil, 552.
 Alabaster, 112.
 Alant camphor, 555.
 Albumen, 535.
 „ detection of, in urine, 591.
 „ vegetable, 539.
Albumen ovi, 535.
 Albumenoid substances, 507, 535.
 Alchemy, 2, 214.
 Alcohol, absolute, 505, 615, 629.
 „ amylic, 525.
 „ and allied bodies, 501 *et seq.*
 „ benzylic, 566.
 „ butylic, 424.
 „ cinnamic, 566.
 „ ethylic, 501.
 „ from sugar, 501.
 „ glyceric, 533.
 „ in 100 parts of spirits of different densities, Table showing the proportion of, 576 in 2nd edition.
 „ methylic, 517.
 „ phenic, 528.
 „ quantitative estimation of, 728.
 „ radicals, 516.
 „ real, 505.
 „ test for impurities in, 736.
 „ test for purity of, 506, 662.
Alcohol amylicum, 525.
 „ *amylicum*, impurities in, 736.
 Alcoholates of chloral, 524.
 Alcoholic drinks, 503.
 Alcoholometer, 631.
 Alcohols, 501.
 Aldehyd, 506.
 „ benzoic, 396.
 „ cinnamic, 554.
 „ euodic, 556.
 „ lauric, 556.
 „ rutic, 556.
 Aldehyds, 515.
 „ aromatic, 551.
 Ale, 478, 508.
 Alexandrian senna, 493.
 Algaroth's powder, 282.
 Alisarin, 569.
 Alkalies, analytical separation of the, 107.
 „ antidotes to, 305.
 „ quantitative estimation of the, 646.
 Alkalimetry, 654.
 Alkaline carbonates, volumetric estimation of the, 651.
 „ earths, 137.
 „ solution of arsenic, 185.
 Alkaloids, 445.
 „ antidotes to the, 448.
 „ nomenclature of, 448.
 „ poisonous, examination for, 586 *et seq.*
 „ reagents for, 588.
 Alkanet, 569.
Alkanna tinctoria, 569.
 Allotropes, 431.
 Allotropic bodies, 481.
 Allotropy, 480.
 Alloxan, 422.
 Alloy, 221.
 Alloys, analysis of, 439.
 Allyl, 532.
 „ cyanide, 533.
 „ sulphide, 532.
 „ sulphocyanate, 532, 556.
 Almond-oil, 547.
 Almonds, oil of bitter, 491, 552.
 „ oil of bitter, test for nitrobenzol in, 530.
Aloe barbadensis, 531.
 „ *socotrina*, 531.
 Aloes, 531.
 Aloins, 531.
 „ formulæ of, 532.
Alstonia scholaris, 467.
 Alstonicine, 467.
 Alstonine, 467.
Athæa officinalis, 454.
 Alum, 150.
 „ cake, 151.
 „ chrome-, 151, 272.
 „ dried, 151.
 „ flour, 151.
 „ impurities in, 736.
 „ iron-, 151.
 „ potash-, 151.
 „ roche or rock, 152.
 „ root, 420.
 „ soda-, 151.
Alumen, 150.
 „ *essiccatum*, 151.

Alumina, 151.

Aluminium, 149.

- „ analytical reactions of, 152.
- „ and ammonium, sulphate of, 150.
- „ and sodium, double chloride of, 150.
- „ bronze, 150.
- „ derivation of word, 24.
- „ detection of, in presence of iron and zinc, 180.
- „ hydrate, 152.
- „ oxide, 151.
- „ quantitative estimation of, 685.
- „ separation of, from chromium and iron, 272.
- „ silicate, 151.
- „ sulphate, 151.

Amalgam, 221.

„ ammonium, 95.

Amber, 415.

„ oil of, 415.

American turpentine, 557.

Amianth, 413.

Amide, 233.

Amido-chloride of mercury, 232.

Amidogen, 232.

Amines, 446.

Ammonia, 94, 96.

- „ detected by Nessler test, 648.
- „ fetid spirit of, 99.
- „ gas, composition of, 96, 141.
- „ gas in solutions of ammonia of different densities, Table showing the amount of, 574 in 2nd edit.
- „ in drinking water, 648.
- „ preparation of, 96.
- „ solution of, 96.
- „ solution of sulphide of, 101.
- „ type, 423.
- „ volcanic, 95.
- „ volumetric estimation of solutions of, 647.

Ammoniacal liquor, 94.

„ salts, sources of, 94.

Ammoniacum, 564.

Ammonia acetatis liquor, 97.

„ aromaticus, spiritus, 98.

Ammonia benzoas, 100, 398.

- „ benzoas, impurities in, 786.
- „ carbonas, 97.
- „ carbonas, impurities in, 786.
- „ citratis, liquor, 99.
- „ fetidus, spiritus, 99.
- „ fortior, liquor, 96.
- „ liquor, 96.
- „ nitras, 99.
- „ nitras, impurities in, 736.
- „ phosphas, 100.
- „ sulphas, 95.

Ammoniated mercury, 232.

„ „ varieties of, 233.

Ammonii bromidum, 100, 312.

„ chloridum, 94.

„ chloridum, impurities in, 736.

Ammonio-chloride of mercury, 232.

„ -citrate of iron, 169.

„ -magnesian phosphate, 181, 386.

„ -nitrate of silver, 198, 234.

„ -sulphate of copper, 198.

„ -sulphate of magnesium, 705.

„ -tartrate of iron, 171.

Ammonium, 94.

- „ acetate, 97.
- „ -amalgam, 95.
- „ analytical reactions of, 104.
- „ and magnesium, arseniate, 181.
- „ and magnesium, phosphate, 131.
- „ and platinum double chloride, 105, 284.
- „ arseniate, 186.
- „ aspartate, 407.
- „ benzoate, 99, 375.
- „ bicarbonate, 98.
- „ bromide, 100, 312.
- „ carbamate, 98.
- „ carbonate, 97.
- „ carbonate, solution of, 99.
- „ chloride, 94.
- „ citrate, 99.
- „ cyanate, 395.
- „ derivation of word, 24.
- „ derivatives, 234.
- „ hydrate, 96.
- „ iodides, 313, 315.

- Ammonium**, molybdate, 387.
 " nitrate, 99, 334.
 " oxalate, 100.
 " peroxide, 315.
 " phosphate, 99.
 " potassium and sodium, separation, of, 107.
 " quantitative estimation of, 679.
 " salts, source of, 94.
 " salts, volatility of, 105.
 " sulphate, 95.
 " sulphide, 101.
 " sulphhydrate, 101.
 " tartrate, 105.
 " volumetric estimation of carbonate of, 649.
- Amomum melegueta*, 555.
- Amorphous**, alkaloid, 459.
 " meaning of, 207.
 " phosphorus, 383.
- Amygdala amara*, 490, 548.
 " *dulcis*, 490, 548.
- Amygdalin**, 490.
- Amyl**, 516.
 " acetate, 525.
 " nitrite, 526.
 " valerianate, 526.
- Amyl nitris*, 526.
- Amylaceous substances**, 473.
- Amylamine**, 447.
- Amylic alcohol**, 525.
- Amylum*, 473.
 " impurities in, 736.
- Amyric acid**, 564.
- Amyrin**, 564.
- Anacyclus pyrethrum*, 562.
- Analogies between chlorine, bromine, and iodine**, 315, 321.
- Analogy of carbon, boron, and silicon**, 415.
 " of oxygen, sulphur, selenium, and tellurium, 352.
 " of sodium and potassium salts, 91.
- Analysis**, 62, 79, 106, 114.
 " blowpipe, 146.
 " gas-, 421, 576.
 " gravimetric, 609, 678.
 " meaning of word, 62.
 " of gases and vapours, 576.
 " of insoluble salts, 433 *et seq.*
 " of medicines, 578.
 " of salts, 433.
- Analysis of substances having unknown properties**, 573.
 " organic, 709.
 " practical, 106.
 " proximate, 709.
 " qualitative, 79, 106, 433.
 " quantitative, 608, 643.
 " spectrum-, 576.
 " systematic, for the detection and separation of the metals, 107, 135, 180, 209, 213, 251, 253, 298.
 " ultimate, 709.
 " volumetric, 609.
- Analytical chemists**, 2.
 " detection of the acidulous radicals of salts soluble in water, 426.
 " memoranda, 255, 298.
- Analytical reactions of**
 acetates, 349.
 albumen, 535.
 alcohol, 506.
 aldehyd, 507.
 aluminium, 152.
 ammonium, 104.
 amygdalin, 490.
 antimony, 206, 579.
 arsenicum, 190, 579.
 atropia, 463.
 barium, 110.
 beberia, 465.
 benzoates, 393.
 berberia, 465.
 bile, 545.
 bismuth, 291.
 borates, 391.
 bromides, 313.
 brucia, 461.
 cadmium, 286.
 caffenia, 471.
 calcium, 133.
 carbonates, 366.
 chloral hydrate, 534.
 chlorates, 342.
 chlorides, 310.
 chromates, 274.
 chromium, 274.
 citrates, 380.
 cobalt, 265.
 conia, 467.
 copper, 216.
 cyanides, 336.
 ferric salts, 178.
 ferridcyanides, 400.
 ferrocyanides, 398.
 ferrous salts, 176.
 fluorides, 400.

Analytical reactions of

formates, 396.
gallic acid, 421.
glucosides, 490 *et seq.*
glycerine, 534.
gold, 282.
guaiacin, 406.
hippurates, 397.
hydrocyanic acid, 326, 328.
hypochlorites, 339.
hypophosphites, 403.
hyposulphites, 405.
iodides, 316.
iron, 176.
lactates, 406.
lead, 242.
lithates, 422.
lithium, 259.
magnesium, 130.
malates, 407.
manganese, 268.
meconates, 408.
mercuric salts, 232.
mercurous salts, 235, 226,
231.
metaphosphates, 408.
morphia, 450.
nickel, 268.
nicotia, 469.
nitrates, 334.
nitrites, 409.
nitrous ether, 513.
oxalates, 369.
phosphates, 386.
phosphites, 403.
platinum, 283.
potassium, 79.
pyrogallie acid, 421.
pyrophosphates, 412.
quinia, 455.
salicin, 408.
silicates, 414.
silver, 249.
sodium, 92.
starch, 474.
strontium, 260.
strychnia, 460.
succinates, 416.
sugar, 484.
sulphates, 362.
sulphides, 354.
sulphites, 357.
sulphocyanates, 416.
tannic acid, 418.
tartrates, 375.
theia, 470.
tin, 278.
urates, 422.

Analytical reactions of

veratria, 471.
zinc, 147.
Anamirta cocculus, 498.
Anchusa tinctoria, 569.
Anchusin, 569.
Andrographis paniculata, 472.
Andropogon citratus, 557.
" *nardus*, 554.
" *schænanthus*, 555.
Aneroid barometer, 611.
Anethene, 552.
Anethol, 553.
Angelic acid, 553.
" powder, 202.
Angelica, 565.
Angostura bark, 473.
" " false, 459.
Anhydride, acetic, 350.
" antimonious, 202.
" antimonious, 202.
" boric, 390.
" carbonic, 365.
" chlorochromic, 274.
" chromic, 272.
" nitric, 333.
" nitrous, 335, 336.
" phosphoric, 15, 384.
" silicic, 413.
" sulphuric, 361.
" sulphurous, 355.
Anhydrides, 87, 350.
Anhydrous bodies, 87.
" nitric acid, 333.
" perchloride of iron, 161.
" sulphate of copper, 216.
Aniline, 530.
" colours, 572.
" blue, 572.
" green, 572.
" red, 572.
" yellow, 572.
Animal charcoal, 120.
" charcoal, decolourizing power
of, 120.
" rouge, 394.
Animals and plants, complementary
action of air, 9.
Aniseed-oil, 552.
Anise-fruit, 552.
Annatto, 569.
Anthemidis flores, 472, 553.
Anthemis nobilis, 553.
Anthracene, 531, 569.
Anthracite, 275.
Antichlor, 357.
Antidotes to acids, 305.
" alkalis, 305.

Antidotes to alkaloids, 448.
 " antimony, 208.
 " arsenic, 198.
 " barium, 110.
 " carbolic acid, 539.
 " copper, 218.
 " cyanides, 337.
 " hydrochloric acid, 310.
 " hydrocyanic acid, 327.
 " lead, 244.
 " mercury, 236.
 " nitric acid, 338.
 " oxalic acid, 370.
 " silver, 250.
 " sulphuric acid, 363.
 " tin, 280.
 " zinc, 149.
Antimonial wine, 203.
Antimoniate of potassium, 92.
 " sodium, 92.
Antimonic anhydride, 202.
 " chloride, 202.
 " oxide, 203.
Antimonii chlorid, liquor, 201.
 " oxidum, 203.
 " oxidum, impurities in, 736.
Antimonious anhydride, 202.
 " chloride, 201.
 " oxide, 202.
 " oxychloride, 202.
Antimonium nigrum, 200.
 " nigrum, impurities in, 736.
 " sulphuratum, 204.
 " tartaratum, 203, 372.
 " tartaratum, quantitative estimation of antimony in, 688.
Antimoniuretted hydrogen, 207.
Antimony, 200.
 " analytical reactions of, 206.
 " and arsenic, analytical separation of, 209.
 " and potassium, tartrate of, 203.
 " antidote to, 208.
 " black, 200.
 " butter of, 201.
 " chloride, 201.
 " crude, 200.
 " derivation of word, 25.
 " from arsenic to distinguish, 194.
 " hydride, 207.

Antimony in organic mixtures, detection of, 579.
 " oxide, 202.
 " oxychloride, 202, 206.
 " oxysulphide, 204.
 " pentachloride, 202.
 " potassio-tartrate, 203.
 " quantitative estimation of, 688.
 " solution of chloride of, 201.
 " sulphide, 200, 206.
 " sulphur salts of, 204.
 " sulphurated, 204.
 " tartarated, 203.
Antiseptic, 529, 534, 538.
Antozone, 272, 599.
Apatite, 389.
Apomorphia, 451.
Aporetine, 395.
Apothecaries, 2.
Apparatus, xv., 3.
 " for experiments, xv.
 " for volumetric analysis, 644.
 " list of, xvi.
Apple-essence, 526.
 " -oil, 526.
 " -wine, 503.
Aqua Anethi, 552.
 " *Aurantii Floris*, 552.
 " *Aurantii Floris*, impurities in, 736.
 " *Camphoræ*, 553.
 " *Carui*, 552.
 " *Chloroformi*, 521.
 " *Cinnamomi*, 552.
 " *Creasoti*, 529.
 " *Destillata*, 139.
 " *Destillata*, impurities in, 736.
 " *Feniculi*, 553.
 " *fortis*, 333.
 " *Laurocerasi*, 491.
 " *Mentha Piperita*, 552.
 " " *Viridis*, 552.
 " *Pimentæ*, 552.
 " *regia*, 209, 324.
 " *Rosæ*, 552, 556.
 " *Sambuci*, 552.
Arabic acid, 479.
Arabin, 123.
Arachidic acid, 543.
***Arachis hypogæa*, 543.**
Arachis-oil, 543.
Araroba powder, 394.
***Arbor Dianæ*, 250.**
Arbutin, 430, 492.
Archil, 570.

- Arctostaphylos uva ursi*, 492.
 Are, 621, 622.
Areca catechu, 420.
Areca nuts, 420.
Argal, vide Acid tartrate of potassium.
Argent-ammon-ammonium, nitrate of, 234.
Argentii nitras, 239, 248.
 „ *nitras*, impurities in, 736.
 „ *oxidum*, 248.
 „ *oxidum*, impurities in, 736.
Argentio chloride, sulphide, etc., vide Salts of silver.
Argentiferous galena, 245.
Argentum, 25.
 „ *purificatum*, 248.
 „ *purificatum*, impurities in, 736.
Argol, vide Acid tartrate of potassium.
Armenian bole, 569.
Armoracia radix, 553.
Arnatto, 569.
Arnica radix, 561.
Arnica, 561.
Arnotto, 569.
Aromatic aldehyde, 551.
Arrowroot-starch (fig.), 476.
Arsenate of ammonium, 188.
 „ *barium*, 111, 196.
 „ *calcium*, 198.
 „ *copper*, 197.
 „ *iron*, 159, 189, 199.
 „ *magnesium and ammonium*, 181.
 „ *silver*, 197, 249.
 „ *sodium*, 188.
 „ *sodium*, volumetric estimation of, 661.
 „ *zinc*, 198.
Arsenates, 187.
Arsenic acid, 187.
 „ and arsenical solutions, volumetric estimation of official, 665.
 „ *anhydride*, 187.
 „ in carbonate of potassium, solution of, 185.
 „ in hydrochloric acid, solution of, 186.
 „ odour of, 187.
 „ *white*, 185.
Arsenical ores, 185.
 „ *sulphur*, 196.
Arsenicum, 185, 187.
 „ analytical reactions of, 190.
- Arsenicum**, antidotes to, 165, 196.
 „ and antimony, analytical separation of, 209.
 „ *bromide*, 184.
 „ *chloride*, 184.
 „ derivation of word, 24.
 „ detection of, in metallic copper, 192.
 „ detection of, in organic mixtures, 579.
 „ *Fleitmann's test* for, 194.
 „ from antimony, to distinguish, 194.
 „ *hydride*, 193.
 „ *iodide*, 185.
 „ *Marsh's test* for, 193.
 „ quantitative estimation of, 665, 687.
 „ red native sulphide of, 185.
 „ reduction of arseniates to arsenites, 188.
 „ *Reinach's test* for, 192.
 „ sources of, 185.
 „ *sulphide*, 185, 196.
 „ yellow native sulphide of, 185.
Arsenide of cobalt, 268.
Arsenio-sulphide of iron, 185.
 „ *nickel*, 268.
Arsenious acid, 186.
 „ *anhydride*, 186.
Arsenite of copper, 197.
 „ *potassium*, 186.
 „ *silver*, 197.
 „ *sodium*, 186.
Arsenites, 186.
Arseniuretted hydrogen, 193.
Art of chemistry, 2.
Artemisia, 499.
Artificial alkaloids, 445.
 „ *gastric juice*, 540.
Asafetida, 564.
Asbestos, 413.
Asclepedin, 472.
Asclepias tuberosa, 472.
 „ *tuberosa*, impurities in, 736.
Ash, 108.
 „ *black-*, 91.
 „ *bone-*, 119.
 „ *soda-*, 91.
Ashes, analysis of (mixed solids), 433.
Asparagin, 407.

- Aspartate of ammonium, 407.
 Aspirator, water, 360.
Assafetida, 564.
 -ate, meaning of, 73, 77.
 Atees, 463.
 Atis, 463.
 Atmosphere, carbonic acid in, 365.
 " composition of, 17.
 " nitrogen in, 16.
 " oxygen in, 16.
 Atmospheric pressure, quantitative determination of, 610.
 Atom, definition of, 55.
 " weights, definition of, 57.
 Atomic proportions, 49, 222.
 " theory, 48.
 " weights, 49.
 Atomicity, 54.
 Atoms, 29.
 " quantivalence of, 54.
 " quantivalence of, definition of, 57.
Atropa belladonna, 463.
Atropia, 463.
 " impurities in, 736.
Atropis sulphas, 464.
 " sulphas, impurities in, 736.
 Atropine, 463.
 " tropate of, 464.
 Attar of rose, 553.
Aurantii cortex, 472, 553.
 " fructus, 553.
 Auric chloride, 281.
 " sulphide, 282.
Aurum, 26.
Avenæ farina, 474.
 Avignon grains, 568.
 Avogadro's and Ampère's "law," 50.
Azadiracta indica, 472.
 Bael fruit, 419.
 " mucilage, 480.
 Bahia powder, 394.
 Baking-powder, 486.
 Balance, 619.
 Balloons, coal-gas for, 14.
 " hydrogen for, 14.
 Balm-of-Gilead fir, 564.
 Balsam, Canada, 557, 564.
 " copaiva, 563.
 " Gurjun, 563.
 " of Peru, 566.
 " of Storax, 566.
 " of Tolu, 566.
Balsamodendron myrrha, 565.
 Balsams, 565.
Balsamum Peruvianum, 566.
Balsamum Peruvianum, impurities in, 736.
 Tolutanum, 566.
Baptisia, 472.
 Barbadoes aloes, 531.
 Barbaloin, 531.
 Barberry, 465.
 Baric chloride, nitrate, etc., *vide* Salts of barium.
 Barff's protected iron, 155.
 Barium, 109.
 " acetonitrate, 137.
 " analytical reactions of, 110.
 " and calcium, separation of, from magnesium, 135.
 " antidotes to, 110.
 " arseniate, 111.
 " carbonate, 110.
 " carbonate, native, 110.
 " chloride, 109.
 " chromate, 110.
 " derivation of word, 24.
 " detection of, in presence of calcium and magnesium, 124, 135.
 " hydrate, 110.
 " nitrate, 109.
 " oxalate, 112, 369.
 " peroxide, 110.
 " phosphate, 111, 358.
 " quantitative estimation of, 680.
 " salts, antidote to, 110.
 " silico-fluoride, 111.
 " sulphate, 110.
 " sulphide, 110.
 " sulphite, 358.
 Barley-starch (fig.), 476.
 " sugar, 488.
 Barometer, 610.
 Baryta, 110.
 " -water, 110.
 Basalt, 150.
 Base, meaning of, 302.
 " organic, 445.
 Basic, meaning of, 302.
 Bassorin, 123, 479.
 Bastard saffron, 570.
 Basyious radicals, 133.
 Bath brick, 413.
 Bauxite, 150.
 Bay-rum, 503.
 " -salt, 78.
 Bearberry, 420.
 Beaver, 561.
 Beberia or beberine, 464.

Beberia sulphas, 465.
 " *sulphas*, impurities in, 736.
 Bebirine, 464.
 Beer, 478, 503.
 Beeswax, 546.
 Beetroot, 485.
Bela fructus, 419.
Belladonna folia, 463.
 " *radix*, 463.
 Bell-metal, 275.
 Bend glass tubes, to, 5.
 Benné oil, 548.
 Benzene, 523.
 Bensin, 523.
 Bensine-collas, 523.
 Benzoate of ammonium, 100, 393.
 " of iron, 394.
 Benzoated lard, 546.
 Benzoates, 392.
 " analytical reactions of, 393.
 Benzoic acid, 392.
 " aldehyd, 393.
 " glycocine, 397.
 Benzoin, 392, 565.
Benzoinum, 392, 565.
 Benzol, 523, 534.
 Benzoline, 528.
 Benzoyl hydrate, 393.
 " hydride, 393.
 Benzyl benzoate, 566.
 " cinnamate, 566.
 Benzyl alcohol, 566.
Berberia or *berberine*, 465.
Berberis cortex, 465.
 Bergamot-oil, 553.
 Berlin blue, 569.
 " red, 569.
 Berthollet's laws, 444.
 Beryllium, 744.
 Betel nuts, 420.
 Bi-, the prefix, 73.
 Bibasic, *see* Dibasic.
 Bibirine, 464.
 Bibulous paper, 116.
 Bicarbonate of ammonium, 96.
 " potassium, 71, 374, 380.
 " sodium, 85, 374, 380.
 " sodium, chemically pure, 646.
 Bichromate of potassium, 271.
 Bikh, 463.
 Bile, 546.
 " detection of, in urine, 592.
 " test for presence of, 545.
 Biliary calculi, 603.
 Bish, 463.

Bismuth, 287.
 " and ammonium, solution of citrate of, 291.
 " analytical reactions of, 291.
 " carbonate, 290.
 " citrate, 290.
 " derivation of word, 26.
 " hydrate, 292.
 " iodides, double, 292, 589.
 " -lozenge, 289.
 " nitrate, 287.
 " oxide, 289.
 " quantitative estimation of, 664.
 " -salts, composition of, 289, 290.
 " subcarbonate or oxycarbonate, 290.
 " subnitrates, impurities in, 787.
 " subnitrate or oxynitrate, 288.
 " sulphate, 289.
 " sulphide, 291.
Bismuthi carbonas, 290.
 " carbonas, estimation of bismuth in, 689.
 " carbonas, impurities 736.
 " oxidum, 290.
 " oxidum, impurities in, 737.
 " subnitrates, 288.
 " subnitrates, impurities in, 787.
Bismuthum, 287.
 " purificatum, 288.
 " purificatum, impurities in, 737.
 Bisulphide of carbon, 516.
 Bisulphite of lime, 357.
 Bitter almonds, oil of, 492, 552.
 " cassava, 474.
 " principles, 472.
 " sweet, 470.
 Bittern, 311.
 Bituminous coal, 275.
 Bivalence, 54.
 Bivalent radicals, 54, 66, 133.
Bixa orellana, 569.
 Bixin, 569.
 Black antimony, 200.
 " ash, 91, 364.
 " band, 154.
 " bone-, 120.
 " cherry bark, 491.

- Black colouring-matters**, 571.
 „ dyes, 571.
 „ flux, 187.
 „ hydrate of iron, 172.
 „ ink, 179, 571.
 „ -lead, 22.
 „ oxide of copper, 215.
 „ oxide of iron, 178.
 „ oxide of manganese, 262.
 „ oxide of mercury, 230.
 „ pepper, 469.
Bladder-green, 571.
Blanc de Perle, 289, 572.
Bleaching by chlorine, 20.
 „ -liquor, 122.
 „ -powder, 122.
Blende, 142.
Block tin, 276.
Blood, 537.
 „ detection of, in organic matter, 599.
 „ composition of, 537.
 „ hydrocyanic acid in the, 328.
 „ root, 470.
 „ stains, 599.
Blowpipe-analysis, 436.
 „ -flame, 146.
Blue colouring-matters, 570.
 „ cochosh, 478.
 „ copperas, 157, 215.
 „ flag, 478.
 „ indigo, 337.
 „ ointment, 219.
 „ pill, 219.
 „ Prussian, 398.
 „ stone, 215.
 „ Turnbull's, 177, 400.
 „ vitriol, 157, 215.
 „ "Boiled" oil, 547.
Boiling-point, definition of, 614.
Boiling-points of various substances, 615.
Boldine, 554.
Boldo, 553.
"Bonds" (Frankland), 149.
Bonduc seeds, 473.
Bone-ash, 119.
 „ -black, 120.
 „ -earth, 119, 382.
Bones, composition of, 120, 282.
Boracic acid, 389.
 „ acid as an antiseptic, 538.
 „ anhydride, 390.
Borates, 389.
 „ analytical reactions of, 391.
Borax, 389.
 „ bead, 264.
 „ impurities in, 737.
Borax, volumetric estimation of, 612.
Bordeaux turpentine, 557.
Boric acid, *see* Boracic acid.
Borneène, 557.
Borneo camphor, 558.
Boron, 390.
 „ chloride, 390.
 „ derivation of word, 25.
 „ flame, 391.
 „ fluoride, 390.
Borotartrate of potassium, 390.
Bos taurus, 545.
Boswellia, 565.
Bourdon barometer, 611.
Boyle's law, 50, 635.
Brandy, 503, 507.
Brass, 142.
Brazil powder, 394.
 „ wood, 569.
Bread, 486.
 „ aerated, 486.
 „ -making, 486.
Breidin, 564.
Brezilin, 569.
Bright's disease, 591.
Britannia metal, 200, 238, 275.
British gum, 477.
Bromal, 524.
 „ hydrate, 524.
 „ alcoholates, 524.
Bromate of potassium, 78.
Bromates, 78, 313, 343.
Bromic acid, 78, 343.
Bromide of ammonium, 100, 312.
 „ arsenicum, 184.
 „ ethyl, 514.
 „ iron, 161.
 „ potassium, 78, 312.
 „ potassium, volumetric estimation of, 661.
 „ silver, 249, 313.
 „ sulphur, 354.
Bromides, 311.
 „ analytical reactions of, 313.
 „ quantitative analysis of, 646.
 „ separation of, from chlorides and iodides, 318.
Bromine, 311.
 „ analytical separation of, 313.
 „ derivation of word, 25.
 „ its analogy to chlorine and iodine, 315, 321.
 „ solution of, 318.
 „ volumetric estimation of free, 696.

- Bromum*, 311.
 " impurities in, 737.
Bronze, 275.
 " aluminium, 150.
 " coinage, 214.
Bronzing-powder, 279.
Broom-tops, 470.
Brown colouring-matters, 571.
 " hæmatite, 154, 167.
 " rosin, 560.
 " sugar, 485.
Brucia, 461.
Brucine, 461.
Brunswick green, 197.
Bryoidin, 564.
Buchu folia, 473, 553.
Buchu, oil of, 553.
Buckthorn-green, 571.
 " -juice, 493.
 " Bumping," 324.
Bunsen gas-burners, 13.
 " valve, 700.
Burette, Mohr's, 645.
Burgundy pitch, 563.
Burners, gas-, 6, 13.
Burnett's disinfecting fluid, 144.
Burnt ochre, 569.
 " sugar, 489, 571.
 " umber, 571.
Butea frondosa, 419.
Butter, 537, 547.
 " of antimony, 201.
 " of cacao, 546.
 " of cocoa, 547.
 " of kokum, 547.
 " of orris, 555.
Butyl, 515.
 " chloral, 524.
 " sulphocyanate, 516, 553.
Butylic alcohol, 424.
Butyrate, 424.
Butyric acid, 424, 549.
Buxine, 465.
Buxus sempervirens, 465.
By-products, 255.
Cacao-butter, 546.
Cadmii iodidum, 286.
 " iodidum, impurities in, 737.
Cadmium, 285.
 " analytical reactions of, 286.
 " derivation of word, 26.
 " hydrate, 286.
 " iodide, 286.
 " oxide of, 287.
 " sulphide, 286.
Cæsalpinia bonducella, 473.
Cæsalpinia brasiliensis, 569.
Cæsium, 744.
Caffeine, 470.
Cajuput oil, 554.
Cajuputene, 554.
Cajuputol, 554.
Caking coal, 275.
Calabar bean, 469.
Calamine, 142.
Calamus draco, 561.
Calcic sulphate, phosphate, etc., vide Salts of calcium.
Calcii chloridum, 113.
 " chloridum, impurities in, 737.
Calcined magnesia, 130.
Calcis carbonas præcipitata, 115.
 " carbonas præcipitata, impurities in, 737.
 " chlorata, liquor, 123.
 " hydras, 114.
 " hypophosphis, 402.
 " phosphas, 119.
 " phosphas, impurities in, 737.
 " saccharatus, liquor, 115.
Calcium, 112.
 " analytical reactions of, 123.
 " and barium, separation from magnesium, 135.
 " bisulphite, 357.
 " carbonate, 112, 115.
 " carb. of, prepared, 118.
 " chloride, 112.
 " chromate, 124.
 " citrate, 330.
 " derivation of word, 24.
 " -flame, 124.
 " fluoride, 112.
 " fluoride, in bones, 119.
 " gummate, 123.
 " hydrate, 114.
 " hypochlorite, 121.
 " hypophosphite, 402.
 " hyposulphite, 353.
 " in presence of barium and magnesium, detection of, 135.
 " oxalate, 124, 369.
 " oxide, 114.
 " phosphate, 112, 119, 367.
 " polysulphide, 353.
 " quantitative estimation of, 681.
 " silicate, 112.
 " sulphate, 112, 123, 353.
 " sulphite, 357.
 " superphosphate, 363.
 " tartrate, 375.

- Calc-spar, 112.
 Calouli, urinary, 601.
 " urinary, examination of, 601.
 Caliche, 830.
 Calomel, 228, 235.
 " test for corrosive sublimate in, 228.
 " tests for constituents of, 230.
Calotropis, 478.
Calumbæ radix, 465.
 Calx, 114.
 " impurities in, 737.
 " chlorata, 122.
 " chlorata, impurities in, 737.
Cambogia, 564.
 Camphor-laurel, 558.
 " mixture, 558.
 " oil, 558.
 " water, 558.
Camphora, 558.
 " impurities in, 737.
 " officinarum, 558.
 Camphoretic acid, 558.
 Camphoric acid, 558.
 " peroxide, 551.
 Camphore, 558.
 Camwood, 569.
 Canada balsam, 557, 564.
 Canadian turpentine, 557.
 Candle-flame, composition of, 13.
Cannella alba cortex, 473.
 Cane-sugar, 485.
 Cannabene, 561.
 " hydride of, 561.
 Cannabin, 561.
Cannabis indica, 561.
Cantharides, 528.
 Cantharidic acid, 559.
 Cantharidin, 558.
 Caoutchouc, 567.
 Capacity, unit, 621.
 Capillary, 612.
 Caproate of glyceryl, 547.
 Caproic acid, 547, 549.
 Caproyl, 515.
 Caprylate of glyceryl, 547.
 Caprylic acid, 547, 549.
 Capsaicin, 466.
Capsici fructus, 466, 563.
 Capsicia, 466.
 Capsicin, 563.
 Capsicine, 466.
 Capsicum-fruit, resin of, 561, 563.
 " oil, 563.
 Caramel, 484, 571.
 Caraway-oil, 553.
 Carbamate of ammonium, 98.
 Carbasotic acid, 530.
Carbo animalis, 120.
 " animalis purificatus, 120.
 " anim. purif., impurities in, 737.
 " ligni, 120.
 " ligni, impurities in, 737.
 Carbolic acid, 529.
 " acid, antidote to, 528.
 Carbon, 22.
 " bisulphide, 516.
 " combustion of, 22.
 " derivation of word, 22.
 " quantitative estimation of, in organic compounds, 709 et seq.
 Carbonate of ammonium, 97.
 " ammonium, solution of, 97.
 " barium, 110.
 " bismuth, 290.
 " calcium, 112, 115.
 " calcium, prepared, 118.
 " iron, 157.
 " iron, saccharated, 158.
 " lead, 238, 244.
 " lithium, 258.
 " magnesium, 127.
 " potassium, 61.
 " potassium, acid, 71.
 " sodium, 83, 91.
 " sodium, acid, 85.
 " sodium, chemically pure, 616.
 " sodium, manufacture of, 91, 364.
 " strontium, 259.
 " zinc, 142, 145.
 Carbonates, 364.
 " acidulous radical in, 66, 364.
 " analytical reactions of, 366.
 " gravimetric estimation of, 702.
 " volumetric estimation of alkaline, 649.
 Carbonic acid, 22, 364.
 " acid gas, generation of, 72.
 " acid gas, solubility of, in water, 88.
 " anhydride, 365.
 " oxide, 398.
 Carbonization, 108.
 Cardamom-oil, 554.
 " greater, 555.
 " lesser, 554.

- Carica papaya*, 542.
 Charles's process for the valuation of cinchona barks, 719.
 Carolina jasmine, 468.
 Carmine, 394.
 Carminic acid, 394.
 Carnallite, 61.
 Carrageen moss, 479.
 Carrotin, 569.
 Carthamin, 570.
Carthamus tinctorius, 570.
Carum ajowan, 552.
 Carvene, 554.
 Carvol, 552, 554.
 Caryophyllin, 554.
 Cascarilla-oil, 554.
Cascarilla cortex, 473, 554.
 Casein, 537.
 " vegetable, 474, 539.
 Cassia-oil, 554.
Cassia pulpa, 485.
Castilleja elastica, 567.
 Cast iron, 154.
 Castor, 561.
Castor fiber, 561.
 Castor-oil, 548.
Castoreum, 561.
 Castorin, 561.
 Catechin, 419.
 Catechu, 419.
Catechu pallidum, 419.
 " *pallidum*, impurities in, 737.
 Catechuic acid, 419.
 Cathartic acid, 492.
 Cathartogenic acid, 492.
Caulophyllum, 473.
 Caustic, 248.
 " lime, 114.
 " lunar, 248.
 " potash, 62.
 " soda, 84.
 Cayenne pepper (*see* *Capsicum*-fruit).
 Cedra-oil, 553.
 Celestine, 260.
 Cellulin, 482.
 Cellulose, 482.
 " of starch, 475.
 Celsius's thermometer, 613.
 Cements, 414.
 Centiare, 622.
 Centigrade thermometer, 613.
Cephalis ipecacuanha, 467.
Cera alba, 546.
 " *alba*, impurities in, 737.
 " *flava*, 546.
 " *flava*, impurities in, 737.
 Cerasin, 479.
Cerasus virginiana, 491.
 Cerebrin, 587.
 Ceresine, 546.
Cerevisia fermentum, 501.
Ceris oxalas, 261.
 " *oxalas*, impurities in, 737.
 Cerite, 261.
 Cerium, 261.
 " derivation of word, 26.
 " oxalate, 261.
 Cerolein, 546.
 Cerotic acid, 546, 549.
 Cetaceum, 546.
 " impurities in, 737.
 Cetene, 546.
Cetraria, 394, 477.
 Cetraric acid, 394.
 Cetyl hydrate, 546.
 " palmitate, 546.
 Cevadilla, 471.
 Cevadilline, 471.
 Cevadine, 469, 471.
 Ceylon "moss," 479.
 Chalcedony, 413.
 Chalk, 112.
 " precipitated, 116.
 " prepared, 118.
 " stones, 603.
 Chalybeate water, 154.
 Chameleon, mineral, 264.
 Chamomile-oil, 553.
 Char, 108.
 Charcoal, 22.
 " animal, 120.
 " animal, decolourising power of, 120.
 " wood, 120.
 Charles's law, 50.
Charta sinapis, 532.
 Chartreuse, 503.
 Chaulmúgra, 473.
Chavica officinarum, 470.
 Chavico acid, 470.
 Cheese, 537.
 Chemical action by symbols, illustration of, 85.
 " action, definition of, 29, 55.
 " affinity, 31.
 " combination, laws of, 57.
 " combination by volume, laws of, 50 *et seq.*
 " combination by weight, laws of, 43 *et seq.*
 " combination different from mechanical, 22, 29.
 " compound, 22.
 " compound, definition of, 56.

- Chemical, diagram, 42, 43, 63, 64, 65.
 " equation, 42, 62.
 " force, 81 *et seq.*
 " force, conditions for the manifestation of, 34.
 " force, its relations to heat and electricity, 639.
 " formula, definition of, 56.
 " formulas, 36.
 " notation, 35, 42.
 " philosophy, principles of, 29 *et seq.*
 " preparations of the British Pharmacopoeia, 607.
 " symbol, definition of, 56.
 " symbols, 35.
 " toxicology, 577.
 Chemicals, 3.
 " list of, xvii.
 Chemism, 31.
 Chemist and Druggist, 2.
 Chemistry, art of, 2.
 " definition of, 55.
 " derivation of the word, 14.
 " inorganic, 445.
 " object of, 2.
 " organic, 445.
 " science of, 2.
 Chemists, analytical, 2.
 " manufacturing, 2.
 " pharmaceutical, 2.
 Cherry-laurel water, 491.
 " sugar in, 486.
 " -tree gum, 479.
 " wild black, 491.
 Chestnut-brown, 571.
 Chian turpentine, 557.
 Chicory, 477.
 Chili saltpetre, 330.
Chimaphila umbellata, 492.
 Chinese red, 569.
 " yellow, 568.
 Chinoidin, 717.
Chirata, 410.
 Chiratin, 410.
 Chirato-genin, 410.
 Chloral, 520, 522.
 " alcoholates, 524.
 " butyl, 524.
 " croton, 524.
 " hydrate, 522.
 " hydrous, 522.
Chloral hydras, 522.
 " *hydras*, impurities in, 737.
 Chlorate of potassium, 340.
 " of potassium, preparation of oxygen from, 5.
 Chlorates, 339.
 Chlorates, analytical reactions of, 342.
 Chloric acid, 122, 339.
 Chloride of ammonium, 94.
 " antimony, 201.
 " arsenicum, 184.
 " barium, 109.
 " boron, 390.
 " calcium, 112.
 " calcium, removal of iron from, 113.
 " chromium, 272, 273.
 " gold, 281.
 " iron, 161, 162, 163.
 " lead, 242.
 " lime, 121.
 " magnesium, 126.
 " manganese, 262.
 " mercuric ammonium, 233.
 " mercurous ammonium, 234.
 " mercury, 220.
 " platinum, 283.
 " platinum and ammonium, 105, 284.
 " platinum and lithium, 259.
 " platinum and potassium, 80, 284.
 " silicon, 415.
 " silver, 247.
 " sulphur, 354.
 " tin, 277.
 " tin, solution of, 277.
 " zinc, 143.
 Chlorides, 305.
 " estimation of, 695.
 " separation of, from bromides and iodides, 318.
 " tests for, 310.
 Chlorinated lime, 121.
 " lime, volumetric estimation of, 672.
 " soda, solution of, 90.
 " soda, volumetric estimation of, 651.
 Chlorine, 18, 308.
 " acids, 342.
 " as a disinfectant, 20.
 " bleaching by, 20.
 " collection of, 19.
 " derivation of word, 23.
 " its analogy to bromine and iodine, 315, 321.
 " liquid, 309.
 " preparation of, 18.
 " properties of, 20.

- Chlorine**, relative weight of, 21.
 " solubility in water, 19.
 " the active agent in bleach-
 ing-powder, 122.
 " volumetric estimation of,
 696.
 " -water, 19, 308.
Chlorochromic acid, 274.
Chloroform, 519.
 " impurities in, 737.
 " water, 521.
Chlorophyll, 571, 605.
Chocolate, 547.
Cholic acid, 545.
Cholate of sodium, 545.
Cholesterin, 537, 603.
Chondrin, 540.
Chondrus crispus, 479.
Chromate of barium, 110.
 " calcium, 124.
 " lead, 243.
 " potassium and ammo-
 nium, 111.
 " potassium, standard
 solution of red, 665.
 " silver, 250.
Chromates, 272.
 " analytical reactions of,
 272.
 " of potassium, 111.
Chrome-alum, 151.
 " -ironstone, 271.
 " -red, 243, 569.
 " -yellow, 243, 568.
Chromic acid, 272.
 " anhydride, 272.
 " hydrate, 274.
 " salts, 272.
Chromium, 271.
 " analytical reactions of,
 274.
 " chloride, 272, 273.
 " derivation of word, 26.
 " oxide of, 271.
 " separation of, from
 aluminium and iron,
 272.
 " sulphate, 272.
Chromous salts, 272.
Chromule, 571.
Chrysammic acid, 530.
Chrysarobine, 394.
Chrysophanic acid, 394.
Cicuta virosa, 555.
Cicutine, 467.
Cider, 503.
Cimicifuga racemosa, 473.
Cimicifugin, 473.
Cinchona Calisaya, 453.
 " *officinalis*, 453.
 " *succirubra*, 453.
Cinchona flava cortex, 453.
 " *pallida cortex*, 453.
 " *rubra cortex*, 453.
Cinchonia, 458.
Cinchonidia, 459.
Cinchonicine, 459.
Cinchonidia, 458.
Cinchonidine, 458.
Cinchonine, 458.
Cinchovatin, 459.
Cinchovatine, 459.
Cinnabar, 218, 569.
Cinnamein, 566.
Cinnamene, 566.
Cinnamic acid, 394, 566.
 " alcohol, 566.
 " aldehyd, 554.
Cinnamol, 566.
Cinnamomi cortex, 554.
Cinnamon-oil, 554.
Cinnamyl alcohol, 566.
 " cinnamate, 566.
 " hydride, 554, 566.
Cissampelia, 465.
Cissampeline, 465.
Cissampelos pareira, 465.
Citrate of ammonium, 99.
 " bismuth, 290.
 " calcium, 380.
 " iron and quinine, 170,
 455, 722.
 " lithium, 258.
 " magnesium, 180.
 " nicotia, 469.
 " potassium, 73.
 " potassium, volumetric
 estimation of, 652.
 " quinine, 455.
 " silver, 381.
Citratee, 378.
 " analytical reactions of, 380.
 " volumetric estimation of,
 652.
Citric acid, 373.
 " action of heat on, 379.
 " saturating power of, 380,
 743.
Citron-oil, 553.
Citronella-oil, 554.
Citronellol, 554.
Citro-tartrate of sodium, 90.
Citrus, 553.
Citrus bergamia, 378.
Classification of elements, 108, 133,
 187, 297.

- Clausius's theory, 15.
Claviceps purpurea, 561.
 Clay, 150, 418.
 " ironstone, 154.
 Cloves, oil of, 554.
 Coal, 275.
 " -brasses, 154.
 " -gas, 516.
 " -gas for balloons, 14.
 " products of, 516, 572.
 " -tar colours, 572.
 Cobalt, 266.
 " analytical reactions of, 265.
 " arsenide, 266.
 " blue, 570.
 " derivation of word, 26.
 " -glance, 266.
 " hydrate, 267.
 " oxide, 266, 570.
 " separation of, from nickel, 269.
 " sulphate, 267.
 " sulphide, 267.
 Cobaltic ultramarine, 570.
 Cobalticyanide of potassium, 267.
 " nickel, 269.
 Coca, 466.
 Cocaine, 466.
Cocculus indicus, 498.
Coccus, 394, 569.
 " *Ilidis*, 204.
Cochineal, 394, 569.
 Cocoa, 547.
 " nibs, 546.
 " nut, 547.
 " -nut oil, 547.
 Coccatina, 547.
Cocos nucifera, 547.
 Codamine, 450.
 Codeia, 450, 452.
 Cod-liver oil, 547.
 Coffee, 470.
 Cohesion, 55.
 Coinage, copper, 214, 632.
 " gold, 230, 632.
 " silver, 246, 632.
 Coke, 22.
Colchici cornus, 466, 477.
 " *semina*, 466.
Colchicia, 466.
Colchicin, 466.
Colchicine, 466.
Coloother, 168, 569.
 Collection of gases, 5, 6.
 Collidine, 464.
 Collin, 730.
 Colloidion, 483.
Colloidium, 483.
Colloidium flexile, 483.
 Colloid bodies, 730.
Colocynthis pulpa, 493.
Colocynthin, 493.
 Colopholic acid, 561.
 Colophonic acid, 561.
 " hydrate, 561.
 Colophonine, 561.
 Colophony, 560.
 Colouring-matters, 568.
 Combination, chemical, 29.
 " by volume, 50 *et seq.*
 Combining proportions, 47 *et seq.*, 232, 608.
 Combustible, 12.
 Combustion, 12.
 " analysis for carbon and hydrogen, 709 *et seq.*
 " analysis for nitrogen, 713.
 " definition of, 56.
 " spontaneous, 175.
 " supporters of, 12.
 Composition of atmosphere, 17.
 " bismuth salts, 289.
 " oils and fats, 543.
 Compound, chemical, 22.
 " chemical, definition of, 56.
 " chemical, different from mechanical, 29.
 Compounds, 29.
 " of the elements, 59.
 Conchicine, 457.
 Condensation, 138.
 Condenser, 138.
 Condensing-tub, 138.
 " worm, 138.
 Condy's disinfecting fluids, 79, 264.
 Confections, 488.
 Conhydrine, 467.
 Conia, conine, or conicine, 467.
Conis folia, 467.
 " *fructus*, 467.
Conium maculatum, 467.
 " quinine, 457.
 Constant proportions, law of, 43.
 Constitution of alkaloids, 445.
 " matter, 37.
 " salts, 66, 133, 303, 331, 348.
 Construction of formulae, 42.
 Convolvulin, 493.
 Convolvulinol, 493.
Convolvulus Scammonia, 500.
 Conylia, 467.
Copastiba, 563.
 " impurities in, 737.

- Copaiva, 563.
 " oil, 554.
 Copalvic acid, 563.
 Copal, 561.
 Copper, 214.
 " acetate, 216.
 " ammonio-sulphate, 217.
 " analytical reactions of, 216.
 " antidotes to, 218.
 " arseniate, 197.
 " arsenite, 197.
 " black oxide of, 215.
 " blue, 570.
 " coinage, 214, 632.
 " derivation of word, 25.
 " detection of arsenicum in, 192.
 " ferrocyanide, 217.
 " foil, 215.
 " hydrate, 217.
 " hydride, 408.
 " iodide, 318.
 " in organic mixtures, detection of, 579.
 " melting-point of, 616.
 " metallic, 214.
 " oxide, 215.
 " oxyacetate, 216.
 " pyrites, 214.
 " quantitative estimation of, 688.
 " quantivalence of, 215.
 " subacetate, 216.
 " sulphate, 215.
 " sulphate, anhydrous, 216.
 " sulphide, 216.
 " -zinc couple, 698.
 Copperas, blue, 157, 216.
 " green, 157.
 Coptis-root, 465.
 Coriander-oil, 554.
 Cork-borers, 5.
 Corpse fat, 540.
 Correction of the volume of a gas for pressure, 635.
 " for temperature, 635.
 Corrosive sublimate, 226.
 " test for, in calomel, 228.
 Corydalia, 467.
 Cotarnine, 450.
 Coto-bark, 498.
 Cotoin, 493.
 Cotton-wool, 482.
 Cowbane, 555.
 Cowhage, 276.
 Cows' milk, 537.
 Cranesbill, 420.
 Cream, 537.
 " of tartar, 61, 81, 373.
 Creasol, 529.
 Creasote, 529.
 Creasotum, 529.
 " impurities in, 737.
 Cresol, 529.
 Cresotic acid, 531.
 Cresylic acid, 529, 531.
 Creta, 118.
 " *præparata*, 118.
Crinum asiaticum, 500.
 Crith, 636.
 Crocin, 568.
 Crocus (mineral), 168, 569.
 " (vegetable), 568.
Crocus sativus, 568.
 Croton chloral, 524.
 " oil, 547, 548.
 Crown bark, 453.
 Crucibles, 69.
 Crude antimony, 200.
 " potashes, 61.
 Crum's test for manganese, 266.
 Cryohydrates, 88.
 Cryolite, 440.
 Cryptophanic acid, 591.
 Cryptopia, 450.
 Crystallization, water of, 87.
 Crystalloid bodies, 730.
 Cubeb pepper, 470.
Cubeba, 470.
 Cubebene, 554.
 Cubebin, 470.
 Cubebs, oil of, 554.
 Cubic inches in a gallon, 636.
 " nitre, 330.
 Cuca, *see* Coca.
 Cudbear, 570.
 Cumin, 555.
 Cuminic acid, 555.
 Cuminol, 555.
Cuminum cyminum, 555.
 Cummin, 555.
 Cupel, 694.
 Cupellation, estimation of silver by, 694.
 Cupr-diammon-diammonium, sulphate of, 234.
Cupri sulphas, 215.
 " *sulphas*, impurities in, 738.
 Cupric arsenite, 197.
 " compounds, 215.
 " ferrocyanide, 217.
 " hydrate, 217.
 " oxide, 215.
 " sulphate, 215.

- Capric sulphide, 216.
 Cuprous iodide, 318.
 " oxide, 215, 591.
Cuprum, 25, 214.
 Curarine, 461.
 Curaçoes, 503.
Curcuma longa, 557, 568.
 Curcumin, 569.
 Curd soap, 545.
 Curds, 538.
 " and whey, 486, 538.
 Currant, sugar in, 425.
Cusparia cortex, 473.
 Cusparin, 473.
Cusso, 562.
 Cutch, 419.
 Cyanates, 395.
 Cyanic acid, 395.
 Cyanide of allyl, 533.
 " mercury, 323.
 " nickel, 269.
 " potassium, 323.
 " silver, 249, 326.
 Cyanides, 322.
 " analytical reactions of metallic, 326.
 " antidote to, 327.
 " double, 323.
 " quantitative estimation of, 695.
Cyanogen, 179, 322.
 Cyanurets, *vide* Cyanides.
 Cyder, *see* Cider.
Cydonia vulgaris, 480, 491.
 Cymene, 551, 552, 557.
 Cymol, 552, 555.
 Cypripedin, 473.
 Cystin, 596.

 Dahlia, 477.
 Dalleiochin, 456.
 Dalton's atomic theory, 43.
 " laws, 43, 44.
 Dandelion, 477.
Daphne laureola, 562.
 " *mezereum*, 493, 562, 565.
 Daphnetin, 493.
 Daphnin, 493.
Datura alba, 463.
 Daturia, or daturine, 467.
 Dangleish's bread, 486.
 Davy safety-lamp, 14.
 Davyum, 744.
 Deadly nightshade, 463.
 Decantation, 117.
 Decimal coinage, 622.
 " weights, 620.
 Decipium, 744.

 Decoctions, 605.
 Decolourising power of animal charcoal, 120.
 Decrepitation, 435.
 Definition of an atom, 55.
 " atomic weights, 57.
 " a chemical compound, 56.
 " a chemical equation or diagram, 56.
 " a chemical formula, 56.
 " a chemical symbol, 56.
 " a gas, 57.
 " a liquid, 57.
 " a mixture, 55.
 " a solid, 57.
 " an element, 55, 129.
 " chemical action, 55.
 " chemical force, 55.
 " chemistry, 55.
 " combustion, 56.
 " law of diffusion, 56.
 " molecular weights, 57.
 " molecule, 55.
 " quantivalence of atoms, 57.
 Deflagrating flux, 441.
 Deflagration, 75.
 Deliquescence, 91.
 Delphia, 467.
Delphinium staphysagria, 467.
 Density, 628.
 " of vapours, 635.
 Dentifrices, action of, 276.
 Deodorizers, 20.
 Deodorizing liquid, 144.
 " power of animal charcoal, 120.
 Deposits, urinary, 594.
 Derivations of names of elements, 22 *et seq.*
 Derivatives of ammonium, 234.
 Desiccation, 675, 708.
 Destructive distillation, 189.
 Detonation, 75.
 De Valangin's solution, 186.
 De Vrij's processes for estimating the value of cinchona bark and the purity of commercial quinine, 716, 720.
 Dextrin, 477.
 Dextrine-maltose, 488.
 Dextroracemic acid, 373.
 Dextrose, 485.
 Dextrotartaric acid, 373.
 Dhak tree, 419.
 Dhatura, 467.
 Diabetic urine, 485, 486, 592.

- Diagram, chemical definition of, 63.
 Diagrams, chemical, 36, 58, 63.
 Dialysate, 720.
 Dialysis, 729.
 Diallytic iron, 731.
 Dialyzed iron, 731.
 Diamines, 447.
 Diamond, 22.
 Diaphragms, 71.
 Diastase, 477.
 " action of upon starch, 478.
 Dibasic acids, 304.
 Dibasyous radicals, 304.
Dicentra formosa, 467.
 Didymium, 261, 744.
 Diethylamine, 446.
 Diethylia, 446.
 Diffusate, 729.
 Diffusion of gases, 14.
 " law of, definition of, 14, 56.
 Digitalein, 494.
 Digitalin, 493.
Digitalinum, 493.
 Digitaliretin, 493.
Digitalis folia, 494.
 Digitonin, 494.
 Digitoxin, 494.
 Dill-oil, 552.
 Diluenta, 276.
 Dinitrobenzol, 526.
 Dinitrocellulin, 482.
Diospyros embryopteris, 420.
Dipterocarpus turbinatus, 463.
 " *balsamum*, 563.
 " *levis*, 563.
 Disinfectant, chlorine as a, 20.
 Disinfectants, 20.
 Disinfecting fluid, Burnett's, 144.
 " fluid, carbolic acid, 128.
 " fluid, Condy's, 79, 264.
 " powder, 122.
 Dissociation, 638.
 Distillation, 137.
 " destructive, 139.
 " dry, 139.
 " fractional, 502.
 Dita, 467.
 Ditamine, 467.
 Ditain, 467.
 Dithionic acid, 405.
 Doek, 395.
 Dolomite, 126.
 Donovan's solution, 224.
Dorema ammoniacum, 564.
 Double chloride of aluminium and sodium, 150.
 " cyanides, 323.
 Double salts, 80.
 Dover's powder, 468.
 Draconyl, 566.
 Dragon's blood, 561.
 Dried alum, 151.
 Drops, 626.
 Dry distillation, 189.
 Drying-apparatus, 116, 675, 680.
 " -oils, 547.
 Drying *in vacuo*, 116.
 " precipitates, 116, 675, 680.
Dryobalanops aromatica, 558.
Duboisia myoporoides, 468.
Dulcamarin, 470.
 Dulcamara, 474.
 Dulong and Petit's law, 601.
 Dutch camphor, 558.
 Dyads, 133.
 Dyeing by mordants, 152.
 Dyer's saffron, 570.
 Dynamic electricity, production of, 143.
 Dynamicity, 54.
 Earth, bone-, 119.
 Earthenware, 414.
 Earth-nut oil, 548.
 Earths, alkaline, 137.
 Eau de Cologne, 552.
 Ebonite, 567.
 Ebullition, 324.
Ecbalit fructus, 495.
 Ecoboline, 561.
Echites scholaris, 467.
 Effervescing citrate of magnesia, 180.
 " citro-tartrate of sodium, 90.
 " potash-water, 73.
 " soda-water, 88.
 Efflorescence, 91.
 Egg, white of, 536.
 " oil of, 536.
 " yolk of, 536.
 Elasmeter, 631.
 Elaeoptens, 550.
 Elaterin, 495.
Elaterium, 495.
 " impurities in, 495, 738.
 Elder-flower oil, 556.
 Elecampane, 477, 555.
 Electricity, production of dynamic, 143.
 " related to chemical action, 639.
 Element, definition of, 55.
 Elements, 1, 3, 23.
 " ancient, 433.
 " and their compounds, 59.

- Elements, classification of, according
 to analogy, 108.
 " etymology of names of, 23.
 " metallic, 4.
 " non-metallic, 4.
 " of medical or pharma-
 ceutical interest, 3.
 " of pharmaceutical inter-
 est, 8.
 " symbols of, 35, 56.
 " symbols, atomic values,
 and weights of the, 744.
 " symbols of, and deriva-
 tion of name of, 35 *et*
 seq.
 " quantivalence of, 132.
 Elemi, 563.
 Elm, common, 480.
 " -mucilage, 480.
 " slippery, 480.
 " -tannin, 419.
 Emetia, 467.
 Emetine, 467.
 Empirical formulæ, 447.
 Emplastra, 242, 561.
 Emplastrum cerati saponis, 346.
 " plumbi, 242.
 " plumbi iodidi, 241.
 Emulsin, 491, 533.
 Emulsions, 544, 565.
 Enemas, 605.
 English red, 569.
 " blue, 570.
 Eosin, 658.
 Epsom salt, 126.
 Equation, chemical, definition of, 56.
 Equations, 42, 62.
 Equisetic acid, 379.
 Equivalence, 54.
 Equivalents, 745.
 Erbium, 744.
 Ergot, 561.
 Ergota, 561.
 Ergotin, 561.
 Ergotine, 561.
 Ergotinine, 561.
 Ericolin, 492.
 Erlangen blue, 570.
 Erucic acid, 549.
 Erythoreline, 395.
 Erythroxylon coca, 466.
 Eseria, 469.
 Esculin, *see* Æsculin.
 Essence of aniseed, 552.
 " apple, 526.
 " ginger, 523.
 " greengage, 526.
 " melon, 526.
 Essence of mirbane, 528.
 " mulberry, 526.
 " peppermint, 552.
 " pineapple, 526.
 " quince, 526.
 Essences, 552.
 Essentia anisi, 552.
 " mentha piperita, 552.
 Essential oils, *see* Oils.
 " quantitative determination
 of weight, 618.
 Etching, 401.
 Ethal, 546.
 Ether, 508.
 " nitrous, 511.
 Ethereal salts, 515.
 Ethers, 515.
 Ethiop's mineral, 234.
 Ethyl, 514.
 " acetate, 349, 513.
 " bromide, 514.
 " butyrate, 526.
 " hydrate, 501.
 " hydride, 514.
 " iodide, 514.
 " nitrite, 410, 511.
 " osanthyate, 526.
 " oxide, 508.
 " pelargonate, 526.
 " sebacate, 526.
 " suberate, 526.
 " sulphate, 505.
 " zinc, 514.
 Ethylamine, 446.
 Ethylene, 533.
 Ethylia, 446.
 Ethylic alcohol, 501.
 Ethylsulphuric acid, 505, 510.
 Etymology of names of elements, 23.
 Eucalyptol, 555.
 Eucalyptus globulus, 555.
 Euchlorine, 343.
 Eudiometry, 576.
 Eugenic acid, 554.
 Eugenin, 554.
 Euonymin, 473.
 Euodic aldehyd, 556.
 Euphorbium, 564.
 Euphorbon, 564.
 Euxanthate of magnesium, 568.
 Evaporation, 71, 108, 119.
 " *in vacuo*, 119.
 Everett's yellow salt, 325.
 Examinations of the Pharmaceutical
 Society of Great Britain, 2, *vide*
 Prefatory matter.
 Expansion on diluting solution of
 ammonia, 97.

- Explosions of gas, 12.
 Extract of malt, 479.
 Extracts, 606.
Extractum filicis liquidum, 548.
 " *glycyrrhizæ*, 496.
 " *glycyrrhizæ liquidum*, 496.
 " *Saturni*, 240.
 Face-rouge, 396.
 Faeces, 590.
 Fahrenheit's thermometer, 618.
Farina tritici, 474.
 Fat-acids, 549.
 Fats and oils, composition of, 543.
 Fats, etc., analysis of, 732.
 " to determine the melting-point of, 615.
 Fatty bodies, 548.
Fel bovinum purificatum, 545.
 " *purificatum*, impurities in, 738.
 Felspar, 440.
 Fennel-oil, 555.
Fer réduit, 175.
 Fermentation, 502, 503.
 " acetic, 503.
 " alcoholic, 502.
 " ammoniacal, 503.
 " butyric, 502.
 " by soluble albumenoids, 503.
 " lactic, 502.
 " mannitic, 502.
 " putrefactive, 502.
 " viscous, 502.
 Ferrate of potassium, 155.
Ferri acetatis tinctura, 165.
 " *arsenias*, 159, 189.
 " *arsenias*, impurities of, 738.
 " *carbonas*, 158.
 " *carbonas saccharata*, 158.
 " *carbonas saccharata*, impurities in, 738.
 " *et ammoniæ citras*, 168, 169.
 " *et ammoniæ citras*, impurities in, 738.
 " *et ammoniæ citras*, quantitative estimation of iron in, 686.
 " *et quiniæ citras*, 168, 170, 455.
 " *et quiniæ citras*, impurities in, 738.
 " *et quiniæ citras*, quantitative estimation of quinia in, 722.
 " *iodidum*, 22, 160.
 " *oxidum magneticum*, 172.
 " *oxidum magneticum*, impurities in, 738.
 " *perchloridi*, liquor, 163.
Ferri pernitritis, liquor, 174.
 " *peroxidum humidum*, 165.
 " *peroxidum hydratum*, 166.
 " *peroxidum hydratum*, impurities in, 738.
 " *persulphatis, liquor*, 164, 169.
 " *phosphas*, 159.
 " *phosphas*, impurities in, 738.
 " *potassio-tartras*, 168, 171.
 " *pulvis*, 175.
 " *subcarbonas*, 158, 167.
 " *sulphas*, 156.
 " *sulphas*, impurities in, 738.
 " *sulphas essiccata*, 157.
 " *sulphas granulata*, 156.
 " *sulphas granulata*, impurities in, 738.
Ferrio acetate, 165, 349.
 " aceto-nitrate, 174.
 " benzoate, 294.
 " chloride, 161, 163.
 " hydrate, 165.
 " iodate, 344.
 " nitrate, 173.
 " oxide, 166, 167.
 " oxide separated from phosphates and oxalates, 438.
 " oxyiodate, 344.
 " oxysulphate, 157.
 " peroxyhydrate, 166.
 " phosphate, 337.
 " salts, 161 *et seq.*
 " salts, analytical reactions of, 178.
 " sulphate, 164.
 " sulphocyanate, 179, 416.
Ferridcyanide of potassium, 399.
Ferridcyanides, 399.
Ferridcyanogen, 177, 399.
Ferrocyanide of potassium, 323, 397.
 " zinc, 148.
Ferrocyanides, 398.
Ferrocyanogen, 177, 398.
Ferroso-ferric hydrate, 172.
 " oxide, 173.
Ferrous arseniate, 159, 189.
 " bromide, 161.
 " carbonate, 157.
 " chloride, 162.
 " citrate, 172.
 " hydrate, 177.
 " iodide, 160.
 " phosphate, 159.
 " salts, 156 *et seq.*
 " salts, analytical reactions of, 176.
 " sulphate, 156.
 " sulphide, 23, 160, 177, 178.

- Ferrous tartrate**, 172.
Ferrum, 24, 154.
 " *redactum*, 175.
 " *tartaratum*, 168, 171.
 " *tartaratum*, estimation of iron in, 738.
 " *tartaratum*, impurities in, 738.
Ferulaic acid, 564.
Fibrin, 536.
 " vegetable, 539.
Ficus, 485.
 " *elastica*, 567.
Fig, 485.
Filex mas, 548.
Filicic acid, 548.
Filter, to dry, 680.
Filtering-paper, 116, 674.
Filters, 116, 674.
Filtrate, 135.
Fine gold, 282.
Fire-clay, 227.
Fire-damp, 516.
Fixed oils, 547.
 " and volatile oils, difference between, 547.
Flame, oxidizing, 147.
 " reducing, 147.
 " structure of, 18.
Flare, 546.
Flax seed, 430, 547.
Fleitmunn's test for arsenicum, 194.
Flexible collodion, 438.
Flint, 413.
Flores zinci, 146.
Flour, 474.
Flowers of sulphur, 851.
Fluckiger's method for the determination of morphia in opium, 724.
Fluid magnesia, 128.
Fluoric acid, 401.
Fluoride of boron, 890.
 " calcium, 112.
 " calcium in bones, 119.
 " lithium, 258.
 " silicon, 401.
Fluorides, 400.
Fluorine, 401.
 " derivation of word, 25.
Fluor-spar, 112.
Foeniculi fructus, 555.
Foil, copper, 215.
Food, analysis of, 732.
 " elements of, 539.
 " how disposed of in the bodies of animals, 590.
Force, chemical, 81 *et seq.*
Forge-scales, 173.
Formates, 895.
Formic acid, 395, 549.
Formica rufa, 395.
Formula, chemical, definition of, 56.
 " official, 19.
 " officinal, 19.
Formulae, 36, 133.
 " construction of, 53.
 " empirical, 504, 714.
 " graphic, 149.
 " rational, 504, 714.
 " typical, 443.
Formyl, 520.
Fousel-oil, 525.
Fowler's solution, 187.
Foxglove, 494.
Fractional distillation, 502.
Frankincense, Arabian, 565.
 " common, 564.
Frankland's graphic formulae, 149.
Frazinus Ornus, 489.
Free acids, estimated, 655.
Freezing-mixture, 356.
French chalk, 572.
 " turpentine, 557.
Fruit-essences, 526.
Fuchsine, 572.
Fume-cupboard, 102.
Fuming sulphuric acid, 362.
Funnel-tubes, 10.
"Fur" in water-vessels, 367.
Furniture of a laboratory, xvi.
Fusel-oil, 525.
Fusibility of metals, Table of the, 617.
Fusible white precipitate, 233.
Fusing-points of fats, 615.
Fustic, 568.
Gab tree, 420.
Galactometer, 631.
Galactose, 487.
Galbanum, 565.
Galena, 237.
 " argentiferous, 245.
Galenic preparations of the British Pharmacopoeia, 605.
Galipot, 561.
Gall of the ox, 545.
Galla, 417.
Gallie acid, 420.
Gallium, 744.
Gallon, 626, 636.
Galls, English, 417.
 " Aleppo, 417.
Gall-stones, 603.
Galvanic test for mercury, 235.
Galvanized iron, 142.

- Gambier, 419.
 Gamboge, 564, 737.
 Gambogic acid, 565.
 Garancin, 569.
Garcinia indica, 547.
 " *morella*, 564.
 " oil, 547.
 " *pictoria*, 565.
 " *purpurea*, 547.
 Garden thyme, 555.
 Garlic, essential oil of, 532.
 Gas-analysis, 421, 516.
 " -burners, 6, 13.
 " constitution of a, 378.
 " definition of, 57.
 " for balloons, coal-, 14.
 " illuminating, 516.
 " lamp, 6.
 Gases and vapours, analysis of, 421, 576, 635.
 " collection of, 5, 6.
 " correction of the volume of, 635.
 " for pressure, 635.
 " for temperature, 635.
 " diffusion of, 14.
 " law of solubility of, in water, 89.
 " relation of, to liquids and solids, 86.
 " specific gravity of, 634.
 Gastric juice, 540.
 " juice, artificial, 540.
Gaultheria procumbens, 526.
 Gaultheric acid, 526.
 Gay-Lussac's law, 50.
 Gelatigenous substances, 540.
 Gelatine, 540.
 " -sugar, 545.
 " vegetable, 479.
 Gelsemine, 468.
 Gelseminic acid, 468.
Gelsemium sempervirens, 468.
 Gentian bitter, 495.
Gentiana lutea, 495.
Gentiana radix, 495.
 Gentianic acid, 495.
 Gentiogenin, 495.
 Gentiopicrin, 495.
 Gentisic acid, 496.
 Gentisin, 495.
Geranium maculatum, 420.
 " oil, 555.
 Geraniol, 555.
 German silver, 142, 268.
 Gin, 474.
 Gingelly oil, 548.
 Ginger, essence of, 558.
 " -grass, 555.
 " oil, 557.
- Girdwood and Rogers's method for detecting strychnia, 583.
 Glacial acetic acid, 847.
 " phosphoric acid, 385.
 Glass, 414.
 " liquor, 414.
 " rods, 70.
 " soluble, 414.
 " tubes, to bend, 5.
 " tubes, to cut, 5.
 " tubes, to draw out, 117.
 Globulin, 537.
 Glucinum, 744.
 Glucose, 485.
 Glucosides, 490.
 Glue, 540.
 Gluten and glutin, 474.
 Glyceric alcohol, 533.
 Glycerine, 242, 533.
 " tests for, 534.
 Glycerines, 533.
Glycerinum, 534.
 " *acidicarbolici*, 529, 535.
 " *acidigallici*, 535.
 " *aciditannici*, 418, 535.
 " *amylis*, 535.
 " *boracis*, 390, 535.
 " impurities in, 738.
 Glyceryl, 533.
 " caproate, 547.
 " caprylate, 547.
 " hydrate, 533, 544.
 " laurate, 547.
 " margarate, 547.
 " myristate, 547.
 " oleate, 543.
 " palmitate, 543.
 " ricinoleate, 543.
 " rutate, 543.
 " tristearate, 544.
 Glycocholates, 545.
 Glycocine, 545.
 Glycocoll, 545.
 Glycol, 533.
 Glycols, 533.
 Glycollic acid, 533.
 Glycol, 534.
 Glycyrrhetin, 496.
Glycyrrhiza radix, 489, 496.
Glycyrrhizin, 489, 496.
 Gnoscopine, 450.
 Goa powder, 394.
 Gold, 280.
 " analytical reactions of, 282.
 " coin, 280, 632.
 " derivation of word, 26.
 " earth, 568.
 " fine, 282.

- Gold**, jewellers', 280.
 " leaf, 280.
 " mosaic, 279.
 " ochre, 568.
 " perchloride, 281.
 " sulphide, 282.
 " yellow, 568.
Golden seal, 465.
Gossypium, 482.
Gothite, 167.
Goulard's cerate, 240.
 " extract, 240.
 " water, 240.
Gracillaria, 490.
Graham's dialytic process, 729.
 " law of diffusion, 14, 56.
Grain, 626.
Grains of paradise, 555.
Gramme, 621.
 " relation of, to grains, 622.
Granati radices cortex, 419.
Granite, 150.
Granulated phosphorus, 383.
 " tin, 276.
 " zinc, 10.
Granulose, 475.
Grape-sugar, 372, 484.
Grapes, dried, 372.
 " sugar in, 485.
Graphic formulæ, 149.
Graphite, 22.
Grass oils (8), 555, 557.
Gravel, 545.
Gravimetric quantitative analysis, 609, 678.
Gravitation, 618.
Gravity, 618.
Green chloride of iron, 162.
 " copperas, 157.
 " iodide of iron, 160.
 " iodide of mercury, 221.
 " pigments, 571.
 " sulphate of iron, 157.
 " ultramarine, 571.
 " vitriol, 157.
Greengage essence, 526.
Grey powder, 219.
Griffith's mixture, 158.
Ground-nut oil, 548.
Group tests, 297.
Guaiaci lignum, 496.
 " resins, 496.
Guaiacin, 496.
Guaiacol, 529.
Guaiaconic acid, 496.
Guaiacum, resin of, 496.
Guaiaretic acid, 496.
Guaiaretin, 496.
Guaiaretinic acid, 496.
Guano, 423.
Guarana, 470.
Guilandina bonducella, 473.
Guinea grains, 555.
Gulanacha, 473.
Gum, 123, 479.
 " -acacia, 479, 123.
 " arabic, 479, 123.
 " Benjamin, 392.
 " British, 477.
 " cherry-tree, 479.
 " resins, 564.
 " Tragacanth, 479.
Gummate of calcium, 123.
 " of lead, 123.
Gummic acid, 479.
Gun-cotton, 482.
 " -metal, 275.
Gunpowder, 336.
Gurjun balsam, 563.
Gutta percha, 567.
Gutta, 626.
Gynocardia odorata, 473.
Gypsum, 112.
Hæmatein, 570.
Hæmatin, 570.
Hæmatite, brown, 154, 167.
 " red, 154, 167.
Hæmatoxyli lignum, 419, 570.
Hæmatoxylin, 419, 570.
Half-sovereign, weight of the, 280.
Haloid salts, 333.
Hambro' blue, 570.
Hard soap, 544.
Hardness of water, 367.
Heat, latent, 88, 140.
 " related to chemical action, 639.
 " source of, 6.
Heavy carbonate of magnesium, 128.
 " magnesia, 130.
 " spar, 110.
 " white, 110.
Hectare, 623.
Helenin, 555.
Hellebore, black, 497.
 " green, 463, 497.
 " white, 468.
 " white American, 468.
Helleborein, 497.
Helleborin, 497.
Helleborus niger, 497.
 " viridis, 468, 497.
Hemidesmi radix, 396.
Hemidesmic acid, 396.
Hemlock, 467, 555.
Hæmossed calculi, 603.

- Henbane, 468.
 Henry and Dalton's laws, 89.
 Herapathite, 456.
 Hesperidene, 553.
 Hesperidin, 553.
 Heterologous series, 516.
Hevea (Siphonia) Brasiliensis, 567.
 Hexads, 133.
 Hexyl, 515.
Hibiscus esculentus, 480.
 Hippuric acid, 396, 596, 598.
 Hips, 488.
 Hoffner's blue, 570.
 Holway's smelting process, 214.
 Homologous series, 515, 549.
 Honey, 498.
 " poisonous, 468.
 Honeydew, 488.
 Hop, 469, 564.
 " essential oil of, 564.
Hordeum decortiatum, 474.
 " starch of (fig.), 476.
 Horseradish oil, 553.
 Hubbuck's oxide of zinc, 146.
Humulus lupulus, 469, 564.
Hydrargyri, iodidum rubrum, 223.
 " *iodidum rubrum*, impu-
 rities in, 738.
 " *iodidum viride*, 221.
 " *iodidum viride*, impu-
 rities in, 738.
 " *nitratis*, 224.
 " *nitratus acidus liquor*,
 225.
 " *ozidum flavum*, 230.
 " *ozidum flavum*, impuri-
 ties in, 738.
 " *ozidum rubrum*, 229.
 " *ozidum rubrum*, impuri-
 ties in, 738.
 " *perchloridum*, 226.
 " *subchloridum*, 228.
 " *subchloridum*, impuri-
 ties in, 738.
 " *sulphas*, 225.
 " *sulphas*, impurities in,
 738.
 " *sulphuretum cum sul-*
 phure, 234.
 " *sulphuretum cum sul-*
 phure, impurities in,
 738.
Hydrargyrum, 25, 219.
 " impurities in, 738.
 " *ammoniatum*, 233.
 " *ammoniatum*, impuri-
 ties in, 738.
 " *cum creta*, 219.
Hydrargyrum cum creta, impurities
 in, 738.
Hydrastia, 465.
Hydrastis canadense, 465.
 Hydrate of aluminium, 153.
 " ammonium, 96.
 " barium, 110.
 " bismuth, 292.
 " benzoyl, 893.
 " cadmium, 286.
 " calcium, 114.
 " castyl, 546.
 " chromium, 274.
 " cobalt, 267.
 " copper, 217.
 " glyceryl, 502.
 " manganese, 265.
 " nickel, 269.
 " potassium, 62.
 " sodium, 84.
 " zinc, 147.
 Hydrated peroxide of iron, 166, 167.
 " substances, 87.
 Hydrates, composition of, 66.
 " identified, 441.
 Hydraulic cement, 414.
 Hydric acetate, chloride, nitrate,
 sulphate, etc., *vide* the respective
 acids, acetic, hydrochloric, etc.
 Hydride of antimony, 207.
 " arsenicum, 193.
 " benzoyl, 393.
 " cinnamyl, 554.
 " copper, 403.
 " ethyl, 514.
 " methyl, 516.
 " phosphorus, 402.
 " silicon, 415.
 Hydrides, 133.
 Hydriodic acid, 314.
Hydrium, 9.
 Hydrobromic acid, 311.
 " ether, 514.
 Hydrocarbons, 556.
 Hydrochlorate of morphia, 448.
 Hydrochloric acid, 21, 305.
 " acid, analytical reac-
 tions of, 310.
 " acid, antidote to, 310.
 " acid, common, 306.
 " acid, dilute, 306.
 " acid in organic mix-
 tures, detection of,
 578, 581.
 " acid in the liquid acid
 of different densities,
 Table of the amount
 of, 570, 2nd edit.

- Hydrochloric acid, volumetric estimation of, 657.
 Hydrocotarnine, 460.
Hydrocotyle asiatica, 478.
 Hydrocyanic acid, 322.
 " acid, analytical reactions of, 326.
 " acid, antidotes to, 327.
 " acid, dilute, 324.
 " acid, from bitter almond and cherry-laurel, 491.
 " acid in organic mixtures, detection of, 582.
 " acid in the blood, 328.
 " acid, Schönbein's test for, 328.
 " acid, volumetric estimation of, 659.
 Hydroferridocyanic acid, 399.
 Hydroferrocyanic acid, 397.
 Hydrofluoric acid, 400.
 Hydrogen, 9.
 " antimonirettered, 207.
 " arseniuretted, 193.
 " benzoate, borate, etc., *vide* the respective acids, benzoic, boracic, etc.
 " combustion of, 11.
 " derivation of word, 23.
 " explosion of, 11.
 " functions of, 133.
 " in artificial light-producers, 12.
 " lightness of, 12.
 " peroxide, 110.
 " persulphide, 352.
 " phosphoretted, 402.
 " preparation of, 10.
 " properties of, 11, 14.
 " quantitative estimation of, in organic compounds, 709 *et seq.*
 " salts of, 302.
 " siliciuretted, 415.
 " sulphuretted, 102, 350.
 " type, 443.
 " used for balloons, 14.
 " weight compared with air, 14.
 " weight of 1 litre, 636.
 " weight of 100 cubic inches, 636.
Hydrogenium, 9.
 Hydrokinone, 492.
 Hydrometers, 631.
 Hydrosulphuric acid, 350.
 Hydrosulphyl, 352.
 Hydrous chloral, 522.
 " compounds, 87.
 Hydroxyl, 66, 352.
 Hygiene, 2.
 Hyocyamia, 468.
Hyoscyami folia, 468.
 Hyoscyamine, 468.
 Hyper-, meaning of, 162.
 Hypo-, meaning of, 403.
 Hypobromites, 313.
 Hypochloride of sulphur, 355.
 Hypochlorite of calcium, 122.
 " sodium, 89.
 Hypochlorites, 339.
 Hypochlorous acid, 339.
 Hypophosphoric acid, 412.
 Hypophosphite of calcium, 403.
 " sodium, 403.
 Hypophosphites, 401.
 Hypophosphorous acid, 401.
 Hyposulphite of calcium, 353.
 " sodium, 404.
 " sodium, standard solution of, 670.
 Hyposulphites, 404.
 Hyposulphurous acid, 404.
 -ic, meaning of, 77, 155.
 Icacin, 564.
 Iceland moss, 394, 477.
 -ide, meaning of, 77.
 Igasuria, 461.
 Ignition, 108.
Illicium anisatum, 553.
 Illuminating agents, analysis of, 732.
 Incense, 565.
 Inch, 626.
 Incineration, 108.
 " of filters in quantitative analysis, 674.
 Indelible ink, 248.
 Indian barberry, 465.
 " gamboge, 564.
 " hemp, 561.
 " ink, 571.
 " ipecacuanha, 468.
 " liquorice, 496.
 " mustard, 533.
 " pennywort, 473.
 " red, 569.
 " rubber, 567.
 " rubber, vulcanised, 567.
 " yellow, 568.
 Indican, 337.
 Indicator, 644.
 Indigluuin, 337.

- Indigo, 337.
 " -blue, 337.
 " sulphate of, 337.
 " -white, 337.
 " wild, 473.
 Indigogen, 337.
 Indigotin, 337.
 Indium, 744.
 Infusible white precipitate, 238.
 Infusions, 605.
 Inhalation of chlorine, 20.
 " conine, 467.
 " hydrocyanic acid, 326.
 Inhalations, 605.
Injunctio morphiae hypodermica, 449.
 Ink, black, 179.
 " Heberden's, 179.
 " indelible, 248.
 " Indian, 571.
 " invisible, 268.
 " marking, 248.
 " printer's, 571.
 " sympathetic, 268.
 Inorganic chemistry, 445.
 " compounds, 445.
 Insecticide, 562.
 Introduction, 1.
Inula Helenium, 477, 555.
 Inulic anhydride, 555.
 Inulin, 477.
 Inulol, 555.
 Inverted sugar, 485.
 Invisible ink, 268.
 Iodal, 524.
 Iodate of potassium, 73, 344.
 Iodates, 344.
 Iodic acid, 344.
 Iodide of ammonium, 313.
 " arsenicum, 185.
 " bismuth and potassium, 589.
 " cadmium, 286.
 " copper, 318.
 " ethyl, 514.
 " hydrogen, 315.
 " iron, 22, 160.
 " lead, 241.
 " potassium, 75, 316.
 " silver, 249.
 " starch, 475.
 " sulphur, 316.
 Iodides, 314.
 " analytical reactions of, 316.
 " of mercury, 221, 222, 223.
 " quantitative estimation of, 696.
 " separation of, from bromides and chlorides, 318.
 Iodine, 22, 314.
 " chloride, 315.
 " derivation of word, 23.
 " its analogy to chlorine and bromine, 315, 321.
 " moisture in, 696.
 " solution of, 316.
 " standard solution of, 663.
 " tincture of, 316.
 " volumetric estimation of, 670.
 " -water, 316.
 Iodoform, 506, 521.
 Iodum, 314.
 " impurities in, 788.
Ipecacuanha, 468.
Ipomoea orizabensis, 497.
 " *purga*, 497.
 " *simulans*, 497.
 " *turpethum*, 226.
 Iridin, 473.
 Iridium, 283, 285, 744.
Iris florentina, 555.
 " *versicolor*, 473.
 Irish moss, 479.
 Irisin, 473.
 Iron, 153.
 " acetate, 165, 349.
 " acetonitrate, 174.
 " alum, 151.
 " ammonio-citrate, 168.
 " ammonio-tartrate, 168.
 " analytical reactions of, 176.
 " arseniate, 159, 189, 199.
 " arseniate, volumetric estimation of, 667.
 " black hydrate of, 172.
 " black oxide of, 173.
 " bromide, 161.
 " carbonate, 157.
 " cast, 154.
 " chlorides, 161 *et seq.*
 " citrate, 172.
 " citrate of, and quinine, 168, 170, 455.
 " compounds, nomenclature of, 155.
 " derivation of word, 24.
 " detection of, in presence of aluminium and zinc, 180.
 " galvanized, 142.
 " hydrated peroxide, 166, 167.
 " iodate, 344.
 " iodide, 22, 160.
 " magnetic oxide, 172.
 " magnetic oxide, estimation of iron in, 668.
 " meconate, 408.

- Iron**, nitrate (pernitrate) of, 178.
 „ in official compounds, estimation of, 686.
 „ ore, magnetic, 153.
 „ ore, needle, 167.
 „ ore, spathic, 154.
 „ ore, specular, 154.
 „ oxide, 166, 167.
 „ oxyhydrates, 167.
 „ oxysulphate, 157.
 „ perchloride, 161.
 „ perhydrate, 165.
 „ pernitrate, 173.
 „ peroxide, 166, 167.
 „ peroxyhydrate, 166.
 „ persulphate, 164.
 „ phosphate, 159.
 „ phosphate, volumetric estimation of, 667.
 „ from phosphates and oxalates, separation of peroxide of, 438.
 „ potassio-citrate, 168.
 „ potassio-tartrate, 168.
 „ protected from rust, 155.
 „ pyrites, 154.
 „ quantitative estimation of, 666, 686.
 „ red oxide of, 168.
 „ reduced, 174.
 „ rust of, 155.
 „ saccharated carbonate of, 158.
 „ saccharated carbonate of, volumetric estimation of, 668.
 „ salts, nomenclature of, 155.
 „ scale, compounds of, 168.
 „ separation of, from aluminium and chromium, 272.
 „ sodio-citrate, 168.
 „ sodio-tartrate, 168.
 „ sulphate, 156.
 „ sulphide, 23, 160, 177, 178.
 „ sulphocyanate, 179, 397.
 „ tartrate, 172.
 „ wrought, 154.
Ironstone, clay, 154.
Isinglass, 540.
Isomerides, 480.
 „ physical, 482.
Isomerism, 480.
Isomers, 480.
Isomorphism, the doctrine of, 52.
Isomorphous bodies, 52.
Isonandra gutta, 567.
Ispaghul, 480.
 -ite, meaning of, 77.
- Ivory-black**, 571.
Jaborandi, 469.
Jalap, Mexican male, 497.
 „ resin, 497.
 „ Tampico, 496.
 „ true, 497.
Jalapa, 497.
Jalapæ resina, 497.
 „ resina, impurities in, 738.
Jalapic acid, 497.
Jalapin, 497.
Jalapinol, 497.
Japconitine, 463.
Jaune brilliant, 286.
Jelly, 540.
Jervia, 468.
Jervine, 468.
Juices, 605.
Juniper-oil, 555.
Kainit, 61.
Kaladana resin, 498.
Kalk, 24.
Kalium, 24.
Kamala, 562.
Kariyat, 472.
Kelp, 814.
Kermes mineral, 204.
Kerner's test for the purity of sulphate of quinine, 720.
Kieserit, 126.
Kilbride mineral, 167.
Kiln, 114.
Kilogramme, 622.
Kilolitre, 622.
Kilomètre, 622.
Kinate of quinia, 453.
Kinetic theory, 15.
King's blue, 570.
Kino, 419.
Kinone, 492.
Kiwach, 276.
Kokum butter, 547.
Kola-nut, 470.
Kosin, 562.
Koussin, 562.
Kouasso, 562.
Krameria radix, 419.
Laboratory furniture, xvi.
Lac, 537.
Lac-dye, 579.
Lactates, 406.
Lactic acid, 406, 533.
Lactoglucose, 487.
Lactometer, 506.
Lactose, 486.

- Lactuca*, 473.
Lactucarium, 473.
Lactucin, 473.
Lævogryate, 486.
Lævoræmic acid, 373.
Lævorotation, 484.
Lævotartaric acid, 373.
Lævulose, 485.
Lakes, 132.
Lampblack, 571.
Lamps, gas-, 6.
Lana philosophica, 146.
Lanthanum, 744.
Lanthopine, 450.
Larch bark, 420.
Lard, 546.
 " benzoated, 546.
 " prepared, 546.
Laricis cortex, 420.
Larix europæa, 420, 556.
Larixin, 420.
Larixinic acid, 420.
Latent heat, 88, 140.
Laudanine, 450.
Laudanosine, 450.
Laughing-gas, 99.
Laurate of glyceryl, 547.
Laurel-camphor, 558.
Lauric acid, 549.
 " aldehyd, 556.
Laurocerasi folia, 491.
Lavender-oil, 555.
 " -water, 552.
Lavæsinium, 744.
"Law," Avogadro's and Ampère's, 50.
 " Berthollet's, 444.
 " Boyle's, 50, 635.
 " Charles's, 50.
 " Dalton's, 43, 44.
 " Dulong & Petit's, 639.
 " Gay-Lussac's, 50.
 " Graham's, 14.
 " Henry & Dalton's, 89.
 " Malaguti's, 444.
 " Mariotte's, 50, 635.
Law concerning molecular weight, 53.
Law of constant proportions, 43.
 " diffusion, definition of, 56.
 " multiple proportions, 44, 222.
 " reciprocal proportions, 47.
 " solubility of gases in liquids, 84.
Laws of chemical combination by weight, 43, 57, 226 *et seq.*
 " chemical combination by volume, 50, 57 *et seq.*
- Lead**, 237.
 " acetate, 239.
 " analytical reactions of, 242.
 " antidotes to, 241.
 " carbonate, 244.
 " chloride, 242.
 " chromate, 243.
 " derivation of word, 25.
 " detection of, in organic mixtures, 579.
 " gummate of, 123.
 " hydrato-carbonate, 244.
 " iodide, 241.
 " nitrate, 240.
 " oleate, 242.
 " oxide, 238.
 " oxyacetate, 240.
 " oxychromate, 243.
 " perchloride, 241.
 " plaster, 242.
 " puce-coloured oxide or peroxide of, 240.
 " pyrophorus, 175.
 " quantitative estimation of, 650, 692.
 " quantivalence, 239.
 " red, 238.
 " shot, 238.
 " subacetate of, 239.
 " sugar of, 239.
 " sulphate, 244.
 " sulphide, 243.
 " sulphide, native, 237.
 " test for, in water, 243.
 " tree, 244.
 " volumetric estimation of solutions of acetate of, 650.
 " white, 238.
Leadstone, 153.
Leaf-green, 571.
Lecanora, 570.
Lees, 372.
Legumin, 534.
Lemon-chrome, 243.
 " -juice, 379.
 " -juice, estimation of mineral acids in, 702.
 " -oil, 553.
Length, unit of, 621.
Lentisk tree, 562.
Lepidolite, 258.
Leptandrin, 473.
Levisticum, 565.
Levulose, 485.
Lichen blue, 570.
Lichenin, 477.
Light carbonate of magnesium, 127.
 " carburetted hydrogen, 156.

Light magnesia, 180.
 Lignin, 482.
 Lime, bisulphite, 857.
 " caustic, 114.
 " chloride of, 122.
 " -juice, estimation of mineral acids in, 702.
 " -kiln, 114.
 " -oil, 553.
 " quick, 114.
 " slaked, 115.
 " superphosphate of, 382.
 " -water, 115.
 Limestone, 112.
 " magnesian, 126.
 " mountain-, 126.
 Limonis succus, impurities in, 738.
 Limonite, 167.
 Line, 626.
 Lini farina, 547.
 " semina, 547.
 Liniment of mercury, 219.
 Liniments, 605.
 Linimentum ammoniac, 544.
 " calcis, 544.
 Linoxyn, 548.
 Linseed, 480, 548.
 " -cake, 548.
 " -oil, 547.
 " -tea, 480.
 Linum usitatissimum, 480.
 Liqueurs, 503.
 Liquid, definition of, 57.
 " camphor, 558.
 Liquidambar orientale, 566.
 Liquids, specific gravity of, 628.
 " official specific gravity of, 628.
 Liquor ammonia, 96.
 " ammonia, impurities in, 739.
 " ammonia acetatis, 97.
 " ammonia citratis, 99.
 " ammonia fortior, 96.
 " ammonia fortior, impurities in, 739.
 " antimonii chloridi, 201.
 " antimonii chloridi, estimation of antimony in, 688.
 " antimonii chloridi, impurities in, 739.
 " arsenicalis, 187.
 " arsenicalis, impurities in, 739.
 " arsenici hydrochloricus, 186.
 " arsenici hydrochloricus, impurities in, 739.

Liquor, atropia, 464.
 " atropia sulphatis, 464.
 " bismuthi et ammonia citratis, 291.
 " bismuthi et ammonia citratis, estimation of bismuth in, 690.
 " bismuthi et ammonia citratis, impurities in, 739.
 " calcis, 115.
 " calcis, impurities in, 739.
 " calcis chlorata, 123.
 " calcis chlorata, impurities in, 739.
 " calcis saccharatus, 115.
 " calcis saccharatus, impurities in, 739.
 " chlori, 19, 309.
 " chlori, impurities in, 739.
 " ferri perchloridi, 163.
 " ferri perchloridi fortior, 163.
 " ferri perchloridi fortior, estimation of iron in, 686.
 " ferri perchloridi fortior, impurities in, 739.
 " ferri pernitrat, 174.
 " ferri pernitrat, estimation of iron in, 686.
 " ferri pernitrat, impurities in, 739.
 " ferri persulphatis, 164, 169.
 " ferri persulphatis, estimation of iron in, 686.
 " ferri persulphatis, impurities in, 739.
 " gutta percha, 567.
 " hydrargyri nitratis acidus, 225.
 " hydrargyri nitratis acidus, impurities in, 739.
 " hydrargyri perchloridi, 227.
 " iodi, 316.
 " lithia effervescens, 258.
 " lithia effervescens, impurities in, 739.
 " magnesie carbonatis, 126.
 " magnesie carbonatis, impurities in, 739.
 " magnesie citratis, 130.
 " morphia acetatis, 449.
 " morphia hydrochloratis, 449.
 " plumbi subacetatis, 240.
 " plumbi subacetatis, impurities in, 739.

- Liquor plumbi subacetatis dilutus**, 240.
 „ *potassæ*, 62.
 „ *potassæ*, impurities in, 739.
 „ *potassæ*, neutralizing power of, 678.
 „ *potassæ*, specific gravity of, 630.
 „ *potassæ*, to prepare pure, 67.
 „ *potassæ effervescens*, 73.
 „ *potassæ effervescens*, impurities in, 739.
 „ *potassæ permanganatis*, 263.
 „ *sodæ*, 84.
 „ *sodæ*, impurities in, 739.
 „ *sodæ arseniatis*, 188.
 „ *sodæ chloratæ*, 90.
 „ *sodæ chloratæ*, impurities in, 739.
 „ *sodæ effervescens*, 88.
 „ *sodæ effervescens*, impurities in, 740.
 „ *strychniæ*, 461.
 „ *zinci chloridi*, 145.
Liquorice, 489.
 „ -sugar, 489, 496.
List of apparatus, xv.
 „ chemicals, xvii.
 „ reagents, xvi.
Litharge, 238.
Lithates, 422.
Lithiæ carbonas, 258.
 „ *carbonas*, impurities in, 740.
 „ *citras*, 258.
 „ *citras*, impurities in, 740.
Lithic acid, 422.
Lithium, 258.
 „ analytical reactions of, 259.
 „ and platinum chloride, 259.
 „ carbonate, 258.
 „ citrate, 258.
 „ derivation of word, 25.
 „ flame, 259.
 „ fluoride, 258.
 „ silicate, 258.
 „ sulphate, 259.
 „ urate, 259.
Litmus, 101, 570.
 „ paper, 101.
 „ solution of, 101, 654.
 „ tincture of, 101.
Litre, 621.
 „ relation of, to pints, 622.
Liver of sulphur, 68.
Lixiviation, 91.
Loadstone or lodestone, 153.
Lobelia, 469.
Lobelina, 469.
Lobelina, 469.
Logwood, 401, 570.
 „ solution of, bleached by chlorine, 20.
Long pepper, 469.
Looking-glasses, 276.
Lotio hydrargyri flava, 230.
 „ *hydrargyri nigra*, 231.
Louis-blue, 570.
Loxa bark, 453.
Lozenges, 605.
Lucifers, 15.
Lump-sugar, 435.
Lunar caustic, 248.
Lupulin, 564.
 „ oleo-resin of, 564.
Lupuline, 469.
Lupulinic acid, 564.
Lupulus, 469, 473, 564.
Luteolin, 569.
Luting, 103.
 „ fire-clay, 227.
 „ linseed-meal, 104.
Lycopodium, 548.
Mace, fixed oil of, 548.
 „ vol. oil of, 555.
Madder, 569.
Magenta, 572.
Magnesia, 129.
 „ calcined, 130.
 „ effervescing citrate of, 130.
 „ fluid, 128.
 „ hydrous carbonate of, 129.
 „ impurities in, 740.
Magnesia levis, 130.
 „ *levis*, impurities in, 740.
Magnesia carbonas, 123.
 „ *carbonas*, impurities in, 740.
 „ *carbonas levis*, 127.
 „ *carbonas levis*, impurities in, 740.
 „ *carbonatis, liquor*, 128.
 „ *sulphas*, 126.
 „ *sulphas*, impurities in, 740.
Magnesite, 126.
Magnesium, 126.
 „ analytical reactions of, 130.
 „ euxanthate, 568.
 „ for analytical purposes, 194.

- Magnesium**, and ammonium, arseniate of, 131.
 " and ammonium, phosphate of, 131, 386.
 " and ammonium, sulphate of, 705.
 " carbonate, 127.
 " chloride, 126.
 " citrate, 130.
 " derivation of word, 24.
 " detection of, in presence of barium and calcium, 135.
 " limestone, 126.
 " oxide, 129.
 " phosphate, in bones, 119.
 " purrate, 568.
 " quantitative estimation of, 688.
 " separation from barium and calcium, 131.
 " silicate, 413, 572.
 " sulphate, 126.
Magnetic iron ore, 153.
 " oxide of iron, 172.
Maggie test for mercury, 236.
Maize starch (fig.), 476.
Malachite, 214.
Malaguti's law, 444.
Malate of atropine, 463.
 " nicotine, 469.
Malates, 407.
Male fern-oil, 548.
Malic acid, 407.
Mallow tea, 480.
Malt, 478.
 " extract, 479.
 " substitutes, 488.
Maltose, 478, 485, 487.
 " estimation of, 725.
Manganate of potassium, 78, 263.
Manganese, 262.
 " analytical reactions of, 265.
 " black oxide of, 262.
 " Crum's test for, 266.
 " derivation of word, 26.
 " quantitative analysis of black oxide of, 684.
Manganesii oxidum nigrum, 262.
Manganous chloride, 262.
 " hydrate, 265.
Manganous sulphide, 265.
Mangosteen oil, 547.
Manihot, starch of (fig.), 476.
Manna, 489.
 " impurities in, 740.
 " "Manna," 489.
Mannite, 489.
Manufacturing chemists, 2.
Manures, analysis of, 732.
Maranta, starch of (fig.), 451.
Marschchino, 503.
Marble, 112.
Margarine, 546.
Margosa bark, 473.
Marine soap, 547.
Mariotte's law, 50, 635.
Marking-ink, 248.
Marl, 150.
Marmor album, 112.
Marsh-gas, 516.
Marshmallow, 480.
Marsh's test for arsenicum, 193.
Massicot, 238.
Mastic, 562.
Mastiche, 562.
Mastichic acid, 562.
Masticin, 562.
Maté, 470.
Matice folia, 473.
Matricaria chamomilla, 553.
Mauve, 572.
May-apple, 563.
Meadow-sweet, 499, 526.
 " oil of, 499, 536.
Measures, 619 *et seq.*
Mechanical and chemical combination, difference between, 22, 29.
Mechanical medicines, 276.
Meconate of iron, 408.
 " of morphine, 448.
Meconic acid, 407, 585.
Meconidine, 460.
Meconine, 450.
Meconoisine, 450.
Meerschaaum, 413.
Mel, 488.
 " impurities in, 740.
 " boracis, 390.
 " depuratum, 488.
Melam, 417.
Melasses, 489.
Meleguetta pepper, 555.
Melissa oil, 557.
Melissic acid, 549.
Melissyl, palmitate of, 546.
Mellon, 417.
Melon essence, 526.
Melting-points, Table of, 616.

- Melting-points of fats, etc., to determine, 615.
 " of metals, 617.
 Memoranda, analytical, 255, 298.
Menispermum canadense, 465.
Meniha, 555.
 " *pulegium*, 555.
 Menthene, 555.
 Menthol, 555.
 Mercuric ammonium, chloride of, 233.
 " chloride, 226.
 " cyanide, 223.
 " hexiodide, 315.
 " iodide, 223.
 " nitrate, 224.
 " oxide, 229.
 " oxynitrates, 224.
 " oxysulphate, 226.
 " salts, 220.
 " salts, analytical reactions of, 232.
 " sulphate, 225.
 " sulphide, 234.
Mercurius viva, 202.
 Mercurous ammonium, chloride of, 234.
 " chloride, 223, 235.
 " chromate, 237.
 " compounds, 220.
 " iodide, 221.
 " nitrate, 224.
 " oxide, 230.
 " salts, 220.
 " salts, analytical reactions of, 235.
 " sulphate, 228.
 " sulphide, 234.
Mercury, 218.
 " amido-chloride, 232.
 " ammoniated, 232.
 " ammonio-chloride, 232.
 " analytical reactions of, 231.
 " antidotes to, 236.
 " basic sulphate, 226.
 " black oxide, 230.
 " carbonates, 226.
 " chlorides, 226.
 " derivation of word, 25.
 " detection of, in organic mixtures, 579.
 " formula of, 219.
 " galvanic test for, 235.
 " hexiodide, 315.
 " iodides, 221.
 " magpie test for, 236.
 " native sulphide, 218.
Mercury, nitrates, 224.
 " nomenclature of salts of, 220.
 " of life, 202.
 " oxides, 229.
 " oxynitrates, 224.
 " oxysulphate, 226.
 " oxysulphide, 234.
 " quantitative estimation of, 690.
 " subchloride, 228, 235.
 " sulphates, 225.
 " sulphide, 234.
 " yellow oxide, 230.
 Meta, meaning of, 408.
 Metaboracic acid, 389.
 Metacinnaméin, 566.
 Metagummic acid, 479.
 Metallic elements, 4.
 Metalloids, 4.
Metals, 4.
 " of minor Pharmaceutical importance, 257.
 " quantitative estimation of, 673.
 " Table of the fusibility of, 617.
 Metamerides, 481.
 Metamerism, 481.
 Metantimonic acid, 203.
 Metaphosphates, 408.
 Metaphosphoric acid, 385, 406.
 Metastannates, 278.
 Metastannic acid, 278.
 Metastyrol, 566.
 Metathesis, 79, 114.
 Metavanadates, 389.
 Methyl, 514.
 " conia, 467.
 " hydride of, 516.
 " salicylate of, 526.
 Methyl-protocatechuic aldehyd, 425.
 Methylamine, 447.
 Methylated spirit, 517.
 " sweet spirit of nitre, 518.
 Methylene, dichloride of, 521.
 Methylic alcohol, 517.
 " alcohol detected in presence of ethylic alcohol, 517.
 Mètre, 621.
 " relation of, to inches, 622.
 Metric system, 620 *et seq.*
 Metrical system, weights and measures of, 620 *et seq.*
 Metrical system of weights and measures, its relation to the British, 620 *et seq.*

- Meum, 565.
Mezeri cortex, 493, 562.
 Mesereon, 562.
 Mica, 150.
Mica Panis, 486.
 Microcosmic salt, 486.
 Microscopic examination of urinary sediments, 596.
 Microscopy of starches, 475.
 Microspectroscope, 576.
 Milk, 537.
 " of sulphur, 352.
 " -sugar, 486.
 Mimetesite, 389.
 Mimosic acid, 419.
 Mineral acids, detection of, in organic mixtures, 581.
 " chameleon, 264.
 " kermes, 204.
 " Kilbride, 167.
 " purple, 569.
 " rouge, 168, 569.
 Minerals, general analysis of, 433 *et seq.*
 " special analysis of, 732.
 Minim, 626.
 Minium, 238.
 Mint, 555.
 Mirbane, essence of, 528.
 Mishmi bitter, 465.
Mistura ferri aromatica, 179.
 " *ferri composita*, 158.
 Mixture, different from chemical combination, 22, 29.
 " definition of, 55.
 Mixtures, 605.
 Mohr's burette, 644.
 Moist sugar, 485.
 Molasses, 489.
 Molecular volume, 53.
 " weight, 53, 637.
 " weights, definition of, 57.
 Molecule, definition of, 55.
 Molecules, 81.
 Molybdates, 388.
 Molybdenum, 744.
 " sulphide, 388.
 Monads, 133.
 Monamines, 447.
 Monobasic acids, 304.
 Monobasylous radicals, 304.
 Monobrom-camphor, 558.
 Mononitrocellulin, 482.
 Mordants, 152.
Mori succus, 569.
 Morphia, 448.
 " acetate, 449.
 " analytical reactions of, 450.
 Morphia, detection of, in organic mixtures, 535.
Morphia acetat, 449.
 " *hydrochloras*, 448.
 " *hydrochloras*, impurities in, 740.
 Morphine, 448.
 " hydrochlorate, 448.
 " in organic mixtures, detection of, 585.
 " quantitative estimation of, 723.
 Mosaic gold, 279.
 Mosandrum, 744.
Moschus, 540.
 " *moschiferus*, 540.
 "Mother-liquor," 116.
 Motion from heat, 68.
 Mountain-blue, 570.
 " limestone, 126.
 Mucic acid, 489.
 Mucilage of bael, 480.
 " elm-bark, 480.
 " gum acacia, 123.
 " linseed, 480.
 " marshmallow, 480.
 " slippery elm, 480.
 " squill, 500.
 " starch, 474.
 " tragacanth, 123.
Mucilago acacie, 123.
 " *amyl*, 474.
 " *tragacanth*, 123.
Mucuna pruriens, 276.
 Mucus in urine, 600.
 Mudar tree, 473.
 Mulberry-calculus, 603.
 " essence, 536.
 " -juice, 569.
 " sugar in, 486.
 Mulder's process for estimating alcohol, 728.
 Multiple proportions, law of, second law of combination, 44.
 Murexid, 422.
 Musk, 540.
 " -deer, 540.
 Mustard, 532.
 " artificial oil of, 532.
 " essential oil of, 532, 556.
 " fixed oil of, 545.
 " "plaster," 532.
Mycoderma aceti, 346.
Mylatris cichorii, 558.
Myrcia acris, 503.
 Myristate of glyceryl, 546.
 Myristic acid, 549.
 Myristicene, 555.

- M**
Mristicol, 555.
Myristin, 546.
Myronate of potassium, 533.
Myrosin, 533.
Myrozylon Pereira, 566.
 toluifera, 566.
Myrrh, 565.
Myrrha, 565.
Myrrhic acid, 565.
Myrtus communis, 556.

N
Naphthalic acid, 393.
Naphthalin, 393.
Narceia, 450.
Narcotine, 450.
Narthez asafetida, 564.
Nascent state, 33.
Natal aloes, 531.
Nataloin, 531.
Natrium, 24.
Natural philosophy, 37.
Nectandra Rodiei, 464.
Nectandra cortex, 465.
Nectandria, 465.
Needle iron ore, 167.
Negative radicals, 528.
Neptunium, 744.
Neroli oil, 553.
Nessler test, 648.
Neutral chromate, 111.
Neutralization, 100.
Nickel, 268.
 " analytical reactions of, 268.
 " arsenio-sulphide, 268.
 " cobaltcyanide, 269.
 " cyanide, 269.
 " derivation of word, 26.
 " hydrate, 269.
 " separation of, from cobalt, 269.
 " sulphide, 268.
Nicker nuts, 473.
Nicotia, nicotine, nicotina, or nictyia, 469, 561.
Nicotiana tabacum, 469.
Nihilum album, 146.
Nim, 473.
Niobium, 745.
Nitrate of ammonium, 99, 334.
 " argent-ammon-ammonium, 234.
 " barium, 109.
 " bismuth, 287.
 " iron, 173.
 " lead, 240.
 " mercury, 224.
 " potassium, 74, 330.
 " silver, 247.

Nitrate of silver, standard solution of, 659.
 " sodium, 83, 330.
 " strontium, 260.
Nitrates, 330.
 " analytical reactions of, 334.
 " quantitative estimation of, 697 *et seq.*
Nitre, 330.
 " cubic, 330.
 " sweet spirit of, 410, 511.
Nitric acid, 329, 332.
 " acid, antidotes to, 338.
 " acid in organic mixtures, detection of, 578, 581.
 " acid, volumetric estimation of, 657.
 " anhydride, 333.
 " anhydride, radical, and acid, in the liq. acid of different densities, Table showing the per cent. of, 568, 2nd edit.
 " oxide, preparation of, 334.
 " peroxide, 334.
Nitrite of amyl, 526.
 " ethyl, 410, 511.
 " potassium, 409.
Nitrites, 409.
 " analytical reactions, 409.
 " in water, test for, 409.
Nitrobenzol, 492, 526.
 " in oil of bitter almonds, test for, 492, 530.
Nitrocellulin, 482.
Nitrogen, 16.
 " derivation of word, 23.
 " in the atmosphere, 16.
 " oxides, 336.
 " peroxide, 334.
 " preparation of, 16.
 " properties of, 17.
 " quantitative estimation of, in organic compounds, 713 *et seq.*
 " relative weight of, 17.
Nitrohydrochloric acid, 209, 334.
Nitrous acid, 335, 336, 401.
 " anhydride, 335, 336.
 " ether, 511.
 " oxide, 99, 336.
Nomenclature of salts :
 alkaloids, 448.
 anhydrides, 87.
 anhydrous bodies, 87.
 -ate, 73, 77.
 carbonisation, evaporation, ignition, incineration, 108.

Nomenclature of salts :

- double salts, 80.
- hydrates, 65.
- hydrous bodies, 87.
- ic, -ous, 77.
- ide, -ite, 77.
- iron salts, 155.
- mercury compounds, 220.

Non-drying oils, 547.

Non-metallic elements, 4.

Non-metals, 4.

Nordhausen sulphuric acid, 362.

Norwegium, 745.

Notation, 80, 42.

Notes, analytical, 255, 298, 426.

Nutmeg, expressed oil of, 546.

oil of, 555.

Nutrition, plastic elements of, 539.

Nux vomica, 459.

Oatmeal, 474.

Occlusion, 285.

Ochre, 568.

Octahedron, 191.

CEnanthylic acid, 549.

Official formula, 19.

liquids, specific gravity of, 629.

substances, volumetric estimation of, 647 *et seq.*

Official formula, 19.

Oil, ajowan, 552.

ajwain, 552.

almond, 547.

amber, 415.

aniseed, 552.

apple, 526.

arachis, 548.

benné, 548.

bergamot, 553.

bitter-almond, 491, 552.

bitter-almond, artificial, 528.

boldo, 553.

buchu, 554.

cacao, 546.

-cake, 548.

cajuput, 554.

camphor, 558.

Cannabis indica, 561.

capsicum, 563.

caraway, 553.

cardamoms, 553.

cascarilla, 554.

cassia, 554.

castor, 547, 548.

cedra, 553.

chamomile, 553.

cinnamon, 554.

Oil, citron, 553.

citronella, 554.

cloves, 554.

cocoa-nut, 547.

cod-liver, 547.

copaiva, 554.

coriander, 554.

croton, 547, 548.

cubeb, 554.

cummin, 555.

dill, 552.

earth-nut, 548.

eggs, 536.

elder-flower, 556.

fennel, 555.

garcinia, 547.

garlic, 532.

geranium, 555.

gingelly, 543.

ginger, 557.

ginger-grass, 555.

grains-of-paradise, 555.

ground-nut, 548.

hop, 564.

horseradish, 553.

Indian hemp, 561.

jaborandi, 469.

juniper, 555.

lavender, 555.

lemon, 553.

lemon-grass, 557.

lime, 553.

linseed, 547.

lycopodium, 548.

mace, fixed, 546.

mace, volatile, 555.

male-fern, 548.

mangosteen, 547.

meadow-sweet, 499, 526.

melissa, 557.

mustard, artificial, 532.

mustard, essential, 532, 556.

mustard, fixed, 548.

myrtle, 556.

neroli, 558.

nutmeg, fixed, 546.

nutmeg, volatile, 555.

of vitriol, 360.

olibanum, 565.

olive, 543, 548.

omum, 552.

orange-flower, 553.

orange-rind, 553.

orris, 555.

pennyroyal, 535.

pepper, 469.

peppermint, 555.

petit-grain, 553.

- Oil, *pilocarpus*, 460.
 „ *pimento*, 553.
 „ *ptychotis*, 552.
 „ *rose*, 555.
 „ *rosemary*, 556.
 „ *ruce*, 556.
 „ *saffron*, 568.
 „ *santal-wood*, 556.
 „ *sassafras*, 556.
 „ *savin*, 556.
 „ *sesamé*, 543.
 „ *shark-liver*, 548.
 „ *spearmint*, 555.
 „ *sperm*, 546.
 „ *star-anise*, 553.
 „ *sweet-flag*, 556.
 „ *teal*, 548.
 „ *theobroma*, 546.
 „ *thyme*, 557.
 „ *turmeric*, 557.
 „ *turpentine*, 556.
 „ *valerian*, 557.
 „ *verbena*, 557.
 „ *water-hemlock*, 554.
 „ *winter-green*, 526.
 „ *wood*, 563.
 Oils, analysis, 732.
 „ and fats, composition of, 543.
 „ drying, 547.
 „ essential, 550.
 „ essential, tested for alcohol, 552.
 „ fixed, 547.
 „ non-drying, 547.
 „ volatile, 547, 550.
 „ volatile, process for, 551.
 Ointments, 605.
 Okra, 480.
 Oleate of glyceryl, 543.
 „ lead, 242.
 Oleates, 543.
 Olefant gas, 533.
 Oleic acid, 544.
 Oleine, 543.
 Oleo-resins, 564.
Oleum amygdalæ, 547.
 „ *andropogi citrati*, 557.
 „ *amethi*, 552.
 „ *anisi*, 552.
 „ *anthemidis*, 553.
 „ *arachis*, 543.
 „ *cajuputi*, 554.
 „ *carui*, 554.
 „ *caryophylli*, 554.
 „ *cinnamomi*, 554.
 „ *copaibæ*, 554.
 „ *coriandri*, 554.
 „ *crotonis*, 548.
 „ *cubebæ*, 554.
 „ *juniperi*, 555.
 „ *lavandulæ*, 555.
 „ *limonis*, 553.
 „ *lini*, 547.
 „ *macis*, 546, 557.
 „ *menthas piperitæ*, 555.
 „ *menthas viridis*, 555.
 „ *morruæ*, 547.
 „ *myristicæ*, 515.
 „ *myristicæ expressum*, 543.
 „ *myrti*, 556.
 „ *olivæ*, 543, 548.
 „ *phosphoratum*, 383.
 „ *pimentæ*, 554.
 „ *pulegii*, 520.
 „ *ricini*, 543.
 „ *rosmarini*, 556.
 „ *rutæ*, 556.
 „ *sabina*, 556.
 „ *sinapis*, 556.
 „ *succini*, 415.
 „ *teredinina*, 556.
 „ *theobromæ*, 543.
 Oilbanum, 565.
 Olive-oil, 543, 548.
 Omentum, 546.
 Onum oil, 552.
 Opal, 413.
Ophelia chirata, 410.
 Ophelic acid, 410.
 Opianine, 450.
 Opianic acid, 450.
 Opium, 448.
 „ detection of, in organic mix-
 tures, 586.
 „ estimation of morphia in,
 723.
 „ impurities in, 740.
 Orange-chrome, 243.
 „ -flower, 553.
 „ -flower oil, 553.
 „ -rind oil, 553.
 „ -wine, 508.
 Orchil, 570.
 Orchis tuber, 480.
 Orcin, 570.
 Ordeal-poison, 469.
 Orellin, 569.
 Organic analysis, 709.
 „ bases, 445.
 „ chemistry, 445.
 „ compounds, 445.
 Orpiment, 185, 568.
 Orris, butter of, 555.
 „ oil of, 555.
 Ortho-, meaning of, 406.
 Orthophosphates, 408.
 Orthophosphoric acid, 410.

- Orthovanadates, 389.
Oryza sativa, 474.
 " *sativa*, starch of (fig.), 476.
Os ustum, 119.
 Osmium, 285, 745.
 Otto of rose, 555.
 Ounce, 626.
 Ourari, 461.
 -ous, meaning of, 77, 155.
Ovi vitellus, 536.
 Ox-bile, 545.
 " -gall, 545.
 Oxalate of ammonium, 100.
 " barium, 112, 369.
 " calcium, 124, 369.
 " cerium, 262.
 " silver, 370.
 " sodium, 369.
 " strontium, 261.
 Oxalates, 368.
 " analytical reactions of, 369.
 " from phosphates and ferrio oxide, separation of, 438.
 " quantitative estimation of, 704.
 Oxalic acid, 368.
 " antidotes to, 370.
 " chemically pure, 369.
 " in organic mixtures, detection of, 581.
 " standard solution of, 546.
 Oxide of aluminium, 151.
 " antimony, 202.
 " bismuth, 289.
 " cadmium, 287.
 " calcium, 114.
 " chromium, 271.
 " cobalt, 266, 570.
 " copper, 215.
 " iron, black, 172.
 " lead, 238.
 " magnesium, 129.
 " manganese, 262.
 " mercury, 229.
 " silicon, 415.
 " silver, 248.
 " tin, 277.
 " zinc, 146.
 " zinc, Hubbuck's, 146.
 Oxides of nitrogen, 336.
 " identified, 441.
 Oxidising flame, 147.
 Oxyacetate of copper, 216.
 " lead, 240.
 Oxyacid salts, 332.
 Oxyacids of sulphur, 404.
 Oxybutyric acid, 533.
 Orycarbonate of bismuth, 290.
 Orychloride of antimony, 202, 206.
 Orychromate of lead, 243.
 Oxygen, 4.
 " analogies, 352.
 " derivation of word, 23.
 " from ozone and antozone, 272.
 " in the air, 4, 16.
 " its relation to animal and vegetable life, 9.
 " preparation of, 4.
 " properties of, 8.
 " quantitative estimation of, in organic compounds, 709 *et seq.*
 " solubility in water, 8.
 " specific gravity of, 14.
 " weight of 100 cubic inches, 638.
 Oxygenated water, 110.
 Oxyhydrates of iron, 167.
 Oxyiodate of iron, 344.
 Oxyinel, 498.
 " of squill, 488.
Oxymel Scilla, 488.
 Oxyinitrates of bismuth, 288.
 " mercury, 224.
 Oxyalts, 332.
 Oxysulphate of iron, 157.
 " mercury, 226.
 Oxysulphide of antimony, 204.
 " mercury, 224.
 Ozokerite, 512, 546.
 Ozone, 272, 317, 599.
 Palas tree, 419.
 Palladium, 285, 745.
 Palm-oil, 546.
 Palmitate of cetyl, 546.
 " glyceryl, 546.
 " melissyl, 546.
 Palmitic acid, 549.
 Palmitine, 546.
 Pancreatin, 542.
 Papaine, 542.
Papaver rhoeas, 569.
 " *somniferum*, 448.
 Papaverine, 450.
Papaveris capsula, 448.
 Papaw, 542.
 Paper, bibulous, 116.
 " for filtering, 116.
 Papers, test-, 101.
 Para-, meaning of, 373.
 Paracyanogen, 324, 326.
 Paraguay tea, 470.

- Parallin, 499.
 Paratartaric acid, 873.
Pareira radia, 465.
 Paricine, 465.
 Parietinic acid, 894.
 Parigenin, 499.
 Parilla, 465.
 Paris blue, 570.
 " red, 569.
 Particles, elementary, 80.
 Patent sugar, 488.
 Paul's modification of Kerner's test for
 the purity of sulphate of quinine, 720.
 Pear wine, 503.
 Pearl-barley, 474.
 " -sago starch (fig.), 476.
 " -white, 289.
 Pearlsh, 61.
 Peas, 539.
 Pectin, 479.
 Pelargonic acid, 549.
 Pelargonium, 555.
 Pelletierine, 419.
 Pellitory root, 562.
 Pelosia, 465.
 Pelosine, 465.
 Pennyroyal oil, 555.
 Pentachloride of antimony, 202.
 Pentads, 183.
 Pentathionic acid, 405.
 Pepper, black, 464.
 " cayenne, 444.
 " cubeb, 464.
 " long, 469.
 " oil of, 469.
 " resin of, 469.
 " white, 469.
 Peppermint oil, 555.
 Pepsin, 541.
 " vegetable, 542.
 Peptone, 542.
 Per-, meaning of, 156.
 Perbromates, 313.
 Percha tree, 567.
 Perchlorate of potassium, 342.
 Perchloric acid, 342.
 Perchloride of gold, 281.
 " iron, 161.
 " platinum, 283.
 " tin, 277.
 Perfumes, 553.
 Perhydrate of iron, 165.
 Periodide of ammonium, 315.
 " mercury, 315.
 " potassium, 315.
 Permanganate of potassium, 78, 263.
 " its use in volumetric
 analysis, 663.
 Perry, 508.
 Pernitrate of iron, 173.
 Peroxide of barium, 110.
 " hydrogen, 110, 551.
 " iron, 166, 167.
 " iron, hydrated, 166, 167.
 " lead, 240.
 " nitrogen, 834, 336.
 Peroxyhydrate of iron, 166.
 Persian berries, 568.
 Persulphate of iron, 167.
 Persulphide of hydrogen, 352.
 Peru, balsam of, 566.
 Peruvine, 566.
 Petalite, 258.
 Petit grain oil, 553.
 Petit's method for the determination
 of morphia in opium, 724.
 Petroleum, light, 528.
 " -testing, 732.
 Pettenkofer's test for presence of
 bile, 545.
Peumus boldus, 553.
 Pewter, 200, 238, 276.
 Phæoretine, 395.
 Pharaoh's serpents, 417.
Pharbitis nil, 498.
 Pharbitisin, 498.
 Pharmaceutical Society of Great
 Britain, examinations of, 2.
 Pharmacy, 3.
 Phenic acid, 528.
 " alcohol, 528.
 Phenol, 528.
 Phenolphthalein, 658.
 Phenyl, 528.
 Phenylamine, 530.
 Phillipium, 745.
 Phosphate of ammonium, 89.
 " barium, 111, 388.
 " calcium, 112, 119,
 387.
 " iron, 159, 387.
 " magnesium and am-
 monium, 181, 386.
 " magnesium and am-
 monium from ox-
 alates and ferric
 oxide, separation
 of, 438.
 " magnesium in bones,
 119.
 " silver, 249.
 " sodium, 90.
 " sodium, how pre-
 pared from phos-
 phate of calcium,
 120.

- Phosphates, 332.
 " analytical reactions of, 336.
 " quantitative estimation of, 705.
 Phosphites, 410, 411.
 " test for, 403.
 Phosphomolybdic acid, 337, 533.
 Phosphorated oil, 333.
 Phosphoretted hydrogen, 402.
 Phosphoric acid, 15, 334, 410.
 " acid, diluted, 334.
 " acid, quantitative estimation of free, 705.
 " anhydride, 15.
 Phosphorous acid, 335, 410.
 Phosphorus, 15, 333.
 " acids of, 410.
 " combustion of, 15.
 " derivation of word, 24.
 " detection of, in organic mixtures, 533.
 " granulated, 333.
 " pill, 333.
 " properties of, 15.
 " red or amorphous, 333.
 " trihydride, 402.
 Phthalic acid, 390.
 Phyllocyanin, 571.
 Phylloxanthin, 571.
 Physical isomerides, 480.
 Physics, 37.
Physostigmatis faba, 469.
Physostigmia, 469.
Physostigmine, 469.
 Picric acid, 530.
 Picrotoxin, 493.
 Pigments, 563.
Pigmentum nigrum, 571.
 Pills, 605.
Pilocarpine, 469.
Pilula aloes et ferri, 157.
 " *ferri carbonatis*, 153.
 " *ferri iodidi*, 23.
 " *hydrargyri*, 219.
 " *hydrargyri subchloridi composita*, 229.
 " *phosphori*, 333.
 " *plumbi cum opio*, 239.
 " *quinia*, 455.
 Pimaric acid, 561.
 Pimento oil, 553.
Pimpinella anisum, 553.
 Pine-apple, essence of, 526.
 Pinic acid, 560.
 Pink saucers, 570.
 " the common, 499.
 Pins, 276.
 Pint, 636.
Pinus, 557, 560, 564.
Piper nigrum, 469.
Piperia, 469.
Piperic acid, 470.
Piperidia, 470.
Piperidine, 470.
Piperine, 469.
Pipette, 313.
Pituri, 469.
Pistachia terebinthus, 557.
 Pitch, 564.
 " burgundy, 562.
Pis burgundica, 562.
 " *liquida*, 564.
Plantago ispaghula, 490.
 Plants and animals, complementary action on air, 9.
 Plaster of ammoniacum and mercury, 219.
 " mercury, 219.
 " Paris, 112.
 Plasters, 242, 561.
 Plastic elements of nutrition, 539.
 " sulphur, 251.
 Platinic salts, 233.
 Platinous salts, 233.
 Platinum, 232.
 " analytical reactions of, 233.
 " and ammonium chloride of, 105, 234.
 " and lithium chloride of, 259.
 " and potassium chloride of, 80, 234.
 " and sodium chloride of, 235.
 " black, 234.
 " derivation of word, 26.
 " foil, 232.
 " perchloride, 233.
 " residues, to recover, 235.
 " spongy, 235.
 " sulphide, 233.
 Pleurisy root, 472.
 Plumbago, 22.
Plumbi acetas, 239.
 " *acetas*, impurities in, 740.
 " *carbonas*, 233.
 " *carbonas*, impurities in, 740.
 " *emplastrum*, 242.
 " *iodidum*, 241.
 " *nitras*, 241.
 " *osidum*, 233.

- Plumbi oxidum*, impurities in, 740.
 " *subacetatis, liquor*, 240.
Plumbic acetate, sulphate, etc., *vide*
 Salts of lead.
 " peroxide, 240.
Plumbum, 237.
 "Plummer's pill," 229.
Pocula emetica, 200.
Podophylli radix, 562.
 " *resina*, 562.
 Poisonous alkaloids, 583, 586.
 Poisons, antidotes to, *see* *Antidotes*.
 " detection of, in organic
 mixtures, 577 *et seq.*
 Polybasic acids, 304.
 Polybasylous radicals, 304.
 Polychroite, 568.
Polygala senega, 500.
 Polygalic acid, 500.
 Polymerides, 481.
 Polymerism, 480.
 Polymorphism, 480.
 Polymorphous bodies, 480.
 Polysulphide of calcium, 353.
 Pomegranate-rind, 419.
 " -root bark, 419.
 Porcelain, 414.
 Port wine, 503.
 Porter, 503.
 Portland cement, 414.
 Positive radicals, 523.
 Potash, solution of caustic, 62.
 " solution of caustic, to prepare
 pure, 67.
 " sulphurated, 68.
 " Table of strength of solutions
 of, 575, 2nd edit.
 " vol. estim. of sol. of, 651.
 Potash alum, 151.
 " -water, 78.
 Potashes, 61.
Potassa caustica, 68.
 " *caustica*, impurities in, 740.
 " *sulphurata*, 68.
 " *sulphurata*, impurities in,
 740.
Potassa acetas, 70.
 " *acetas*, impurities in, 740.
 " *bicarbonas*, 71.
 " *bicarbonas*, impurities in, 740.
 " *bichromas*, 271.
 " *carbonas*, 61.
 " *carbonas*, impurities in, 740.
 " *chloras*, 340.
 " *chloras*, impurities in, 740.
 " *citras*, 73.
 " *citras*, impurities in, 740.
 " *effervesens liquor*, 73.
Potassa liquor, 63, 67.
 " *liquor*, to prepare pure, 67.
 " *nitras*, 74, 330.
 " *nitras*, impurities in, 740.
 " *permanganas*, 78, 263.
 " *permanganas*, impurities in,
 740.
 " *prussias flava*, 323, 397.
 " *sulphas*, 74.
 " *sulphas*, impurities in, 741.
 " *sulphurata, unguentum*, 68.
 " *tartras*, 74.
 " *tartras*, impurities in, 741.
 " *tartras acida*, 61, 81, 872.
 " *tartras acida*, impurities in,
 741.
 Potassic hydrate, etc., *vide* *Salts of*
 potassium.
Potassii bromidum, 77.
 " *bromidum*, impurities in, 741.
 " *ferricyanidum*, impurities in,
 741.
 " *iodidum*, 76.
 " *iodidum*, impurities in, 741.
 Potassio-citrate of iron, 168.
 " -tartrate of antimony, 203.
 " -tartrate of iron, 168.
 Potassium, 61.
 " acetate, 69.
 " acid carbonate, *vide*
 Bicarbonate.
 " analytical reactions of,
 79.
 " and platinum, chlo-
 ride, 80, 284.
 " and sodium tartrate,
 89, 374.
 " angelate, 553.
 " antimoniate, 92.
 " bicarbonate, 71.
 " bichromate, 271.
 " borotartrate, 390.
 " bromate, 78.
 " bromide, 77.
 " carbonate, 61, 651.
 " ehlorate, 4, 340.
 " chloride, 79.
 " chromate, 111.
 " citrate, 73.
 " cobalticyanide, 267.
 " cyanate, 395.
 " cyanide, 323.
 " derivation of word, 24.
 " ferrate, 155.
 " ferridcyanide, 399.
 " ferrocyanide, 323, 379.
 " -flame, 61.
 " hydrate, 62.

- Potassium hydrate**, to prepare pure solution, 67.
 " in solutions of potash of different densities, Table showing the amount of oxide of, 575, 2nd edit.
 " iodate, 75, 344.
 " iodide, 75, 316.
 " manganate, 78, 263.
 " myronate, 533.
 " nitrate, 74, 330.
 " nitrite, 409.
 " oleate, 544.
 " perchlorate, 342.
 " periodide, 315.
 " permanganate, 78, 263.
 " preparation of, 61.
 " properties of, 62.
 " quantitative estimation of, 673.
 " quantivalence of, 62.
 " red chromate of, 271.
 " red prussiate of, 399.
 " salts, analogy of, to sodium salts, 91.
 " sodium and ammonium, separation of, 107.
 " sources, 64.
 " sulphate, 74.
 " sulphide, 63.
 " sulphocyanate, 416.
 " tartrate, 74.
 " tartrate acid, 61, 80, 372.
 " triiodide, 315.
 " yellow chromate of, 111.
 " yellow prussiate of, 323, 397.
Potato-oil, 525.
 " -starch (fig.), 473, 476.
Poultices, 605.
Pound, 626.
Powder, bleaching, 122.
Powders, 605.
 " soda-, 90.
 " specific gravity of, 632.
Practical analysis, 166.
Precipitant, 80.
Precipitate, 80.
Precipitated chalk, 116.
 " sulphur, 352.
Precipitates soluble in solutions of salts, 255.
 " to wash, 117.
 " to weigh, 676.
Precipitation, 80.
Preparations of the British Pharmacopoeia, chemical, 607.
Preparations of the British Pharmacopoeia, galenical, 606.
Prepared carbonate of calcium, 118.
 " chalk, 118.
 " lard, 546.
 " suet, 546.
Pressure, correction of vol. of gas for, 635.
 " -ganges, 611.
Prickly ash, 465.
Principles of Chemical Philosophy, 29 *et seq.* (See also "Laws.")
Printer's ink, 571.
Prismatic nitre, 330.
Prollius' method for the determination of morphia in opium, 723.
Proportions, atomic, 43.
 " constant, 43.
 " multiple, 44.
 " reciprocal, 47.
Proof spirit, 504.
Propenyl, 534.
Prophetin, 495.
Propionic acid, 549.
Propyl, 516.
Propylamine, 447.
Protocatechuic aldehyd, 425.
Protopine, 450.
Proximate analysis, 709.
Prune, 488.
Prunum, 488.
Prunus serotina, 491.
Prussian blue, 178, 398, 570.
Prussiate of potash, red, 399.
 " of potash, yellow, 323, 397.
Prussic acid, 322.
Pseudaconitine, 463.
Pseudojervine, 469.
Pseudomorphia, 450.
Pterocarpus lignum, 569.
Pterocarpus santalinus, 569.
Ptyalin, 601.
Ptychotis ajowan, 553.
Puce-coloured oxide of lead, 240.
Puddling iron, 154.
Pulveres effervescentes aperientes, 90, 375.
Pulvis algarothi, 202.
 " angelicus, 202.
 " antimoniatis, 205.
 " specuacanthæ compositus, 468.
Pumice-stone, 413.
Punica granatum, 419.

- Purified ox-bile, 545.
 Purple of Cassius, 232.
 " foxglove, active principle in, 493.
 Purpurine, 595.
 Purrate of magnesium, 568.
 Purree, 568.
 Pus in urine, 600.
 Putty-powder, 278.
Pyrethri radix, 562.
 Pyrethric acid, 562.
 Pyrethrin, 470, 562.
Pyrethrum carneum, 562.
 Pyrites, copper, 214.
 " iron, 184.
 Pyroarsenate of sodium, 188.
 Pyroarsenates, 188.
 Pyrogallie acid, 421.
 " acid, use of, in gas-analysis, 421.
 Pyroligneous acid, 345.
 Pyrolusite, 262.
 Pyrometers, 616.
 Pyromorphite, 389.
 Pyrophorus, 175.
 Pyrophosphates, 412.
 Pyrophosphoric acid, 335, 412.
 Pyrovanadates, 389.
 Pyroxylic spirit, 517.
Pyroxylin, 482.

 Quadrivalence, 54.
 Qualitative analysis, 106, 433.
 Quantitative analysis, 608 *et seq.*
 " determination of atmospheric pressure, 610.
 " determination of temperature, 611.
 " determination of weight, 618.
 Quantivalence, 54, 182.
 " of acidulous radicals, 66.
 " of atoms, definition of, 57.

 Quartz, 413.
Quassia lignum, 473.
 Quassin, 473.
 Quercitrin, 568.
 Quercitron, 568.
Quercus cortex, 418.
 " *tinctoria*, 568.
 Quevenne's iron, 175.
 Quick lime, 114.
 Quinamine, 459.
 Quince seeds, 490, 491.
 Quinia, or quinine, 453.
 " amorphous, 717.

 Quinia, analytical reactions of, 456.
 " citrate, 455.
 " citrate of iron and, 455.
 " De Vrij's process for estimating, 715.
 " disulphate, 454.
 " iodo-sulphate, 456.
 " kinate, 453.
 " quantitative estimation of, 715.
 " sulphate, tests for the purity of, 720.
 " sulphates, 454.
 " wine, 455.
Quinia sulphas, 453.
 " *sulphas*, impurities in, 741.
 Quinicia, 459.
 Quinicine, 459.
 Quinidia, 457.
 Quinidine, 457.
 Quinine, 453, 457.
 Quiniretin, 459.
 Quinoidia, 457, 459.
 Quinivalence, 54.

 Racemic acid, 373.
 Radicals, acidulous, 66, 188.
 " acidulous, formulæ of, 66.
 " alcohol, 515.
 " basylous, 183.
 " definition of, 66.
 " negative, 528.
 " positive, 528.
 Rai, 533.
 Raisins, 372, 462, 488.
 Raspberry, sugar in, 486.
 Ratafia, 503.
 Rational formulæ, 504, 714.
 Reactions, analytical, 62.
 " synthetical, 62.
 Reagents, list of, xvi.
 Real alcohol, 506.
 Realgar, 185.
 Réaumur's thermometer, 613.
 Reciprocal proportions, law of, 47.
 Rectification, 139.
 Rectified oil of turpentine, 557.
 " spirit, 139.
 Red, Chinese, 569.
 " chromate of potassium, 271.
 " chrome, 569.
 " colouring-matters, 569.
 " corpuscles in blood, 537.
 " earth, 569.
 " enamel colours, 570.
 " gravel, 495.

- Red hematite, 154.
 „ iodide of mercury, 223.
 „ lead, 238.
 „ litmus-paper, 101.
 „ ochre, 569.
 „ oxide of iron, 168, 569.
 „ Paris, 564.
 „ phosphorus, 383.
 „ -poppy petals, 569.
 „ precipitate, 229.
 „ prussiate of potash, 399.
 „ -rose petals, 569.
 „ sandal-wood, 569.
 „ sanders-wood, 569.
 „ Venetian, 168.
 Reduced indigo, 337.
 „ iron, 174.
 Reducing flame, 147.
 Reinsch's test for arsenicum, 192.
 Relative weight of hydrogen and oxygen, 14.
 Rennet, 588.
Reseda luteola, 569.
 Resin, 557, 560.
 „ of arnica, 561.
 „ of cannabis, 561.
 „ of capsicum, 561.
 „ of castor, 561.
 „ of ergot, 561.
 „ of guaiacum, 496, 561.
 „ of Indian hemp, 561.
 „ of jalap, 497, 561.
 „ of kamala, 562.
 „ of kousso, 562.
 „ of mastic, 562.
 „ of mezereon, 562.
 „ of pepper, 469, 562.
 „ of podophyllum, 562.
 „ of pyrethrum, 562.
 „ of rattleria, 562.
 „ of scammony, 500.
Resina, 560.
 Resinoid substances, 560.
 Resins, 560.
 Respiratory materials of food, 539.
 Retort, 138.
Rhamni succus, 498.
 Rhamnus, 568.
Rhamnus catharticus, 571.
 „ *infectoriis*, 568.
 Rhaponticin, 395.
 Rhatany root, 419.
Rhei radix, 394.
 „ impurities in, 741.
 Rheic acid, 395.
 Rhein, 395.
 Rheumin, 395.
 Rhodium, 285, 745.
 Rhoadine, 450.
Rhæados petala, 569.
 Rhubarb, oxalate of calcium from, 368, 598.
 Rhubarbaric acid, 395.
 Rhubarbarin, 395.
Rhus cotinus, 568.
 Rice, 474.
 Rice-starch (fig.), 476.
 Ricinine, 548.
 Ricinoleate of glyceryl, 548.
 Ricinoleine, 548.
 Ringworm powder, 394.
Roccella, 570.
 Roche alum, 152.
 Rochelle salt, 74, 89, 374.
 Rock alum, 152.
 „ -salt, 78, 306.
 Rohun-bark, 473.
 Roll sulphur, 351.
 Roman cement, 414.
Rosa canina fructus, 468.
 „ *centifolia petala*, 556, 570.
 „ *gallica petala*, 569.
 Rose-aniline, 572.
 „ -oil, 555.
 „ -petals, 556, 569.
 „ -water, 552.
 Rosemary-oil, 556.
 Rosin, 557, 560.
 Rotang palm, 561.
 Rotten-stone, 150.
Rottlera tinctoria, 562.
 Rottlerin, 562.
 Rouge, animal, 394.
 „ mineral, 168, 569.
 „ vegetable, 569.
Rubia tinctorum, 569.
 Rubian, 569.
 Rubidium, 745.
 Rubijervine, 469.
 Ruby, 150.
 Rue-oil, 556.
 Rum, 508.
 Rumex, 395.
 Rumicin, 395.
 Rust of iron, 155.
 Rutate of glyceryl, 548.
 Ruthenium, 285, 745.
 Rutic acid, 549.
 „ aldehyd, 556.
Sabadilla, 471.
 Sabadilla, or sabadilline, 471.
 Saccharated carbonate of iron, 158.
 „ carbonate of iron, volumetric estimation of, 668.

- Saccharic acid, 484.
 Saccharine, 488.
 " substances, 478.
 Saccharometer, 631, 727.
 Saccharometry, 631, 727.
Saccharomyces cerevisia, 502.
Saccharum lactis, 486.
 " *purificatum*, 485.
 " *ustum*, 489.
 Safety-lamp, 14.
 " -tube, 396.
 Safflower, 570.
 Saffranin, 568.
 Saffron, 568.
 " bastard, 570.
 " dyer's, 570.
 " oil of, 568.
 Saffron, 556.
 Saffrol, 556.
 Sago, 474.
 " starch (fig.), 476.
 Sal-ammoniac, 94.
 Salep, 490.
 Salicin, 498.
 Salicyl, hydride of, 498, 526.
 Salicylate of methyl, 526.
 Salicylic acid, 526, 530.
 Salicylous acid, 526.
 Saligenin, 498.
 Saliretin, 498.
 Saliva, 601.
Sal prunella, 331.
 " volatile, 98.
 Salseparin, 499.
 Salt, common, 83, 306.
 " definition of a, 60.
 " of sorrel, 369.
 Saltpetre, 330.
 " Chili, 330.
 Salts, acid, 352.
 " action of the blowpipe on, 146.
 " action of heat on, 435.
 " action of sulphuric acid on, 436.
 " analogies of, 91.
 " analysis of insoluble, 433.
 " constitution of, 66, 133, 302, 331, 348, 445.
 " constitution of, resumé, 443.
 " formation of, 70.
 " nomenclature of, 77.
 " of ammonium, volatility of, 107.
 " of iron, nomenclature of, 155.
 " physical properties of, 434.
 " substitution of for each other, 91.
 Salts, Table of the solubility or insolubility of, in water, 430.
 Sambucene, 556.
Sambuci flores, 556.
 Sand, 413.
 " -bath, 20.
 " -tray, 20.
 Sandal-wood, oil of, 556.
 " red, 569.
 " white, 556.
 " yellow, 556.
 Sandstone, 413.
Sanguinaria canadensis, 470.
 Santalin, 569.
Santalum album, 556.
 Santonic acid, 499.
 Santonica, 499.
 Santonin, 499.
Santoninum, 499.
 " impurities in, 741.
 Santoniretin, 499.
 Sap-green, 571.
 Sapan-wood, 569.
Sapo animalis, 545.
 " *durus*, 544.
 " *durus*, impurities in, 741.
 " *mollis*, 544.
 " *mollis*, impurities in, 741.
 " *viridis*, 544.
 Saponin, 499.
 Sapphire, 150.
Sarcina ventriculi in urine, 601.
 Sarcocollin, 407.
Sarracenia purpurea, 471.
 Sarsaparilla, 499.
Sarsa radia, 499.
 Sassafras-oil, 556.
Sassafras radia, 556.
 Sassafril, 556.
 Saturated solutions, boiling-points of, 615.
 Saturating power of citric acid, 380, 743.
 " power of tartaric acid, 374, 743.
 Saturation, 69.
 " Tables, 374, 380, 743.
 Saturn, 239.
 Saturnine colic, 239.
 Savin-oil, 556.
 Saxony blue, 570.
 Saxony blue, 570.
 Scale compounds of iron, 168.
Scammonia radia, 500.
 " *resina*, 500.
 " *resina*, impurities in, 741.
 Scammonin, 500.

- Scammoniol, 500.
Scammonium, 500, 741.
 Scammony, resin of, 500.
 Scandium, 745.
 Scents, 552.
 Scheele's green, 197.
 Schist, 150.
 Schönbein's test for hydrocyanic acid, 828.
 Schweinfurth green, 197.
 Science of chemistry, 2.
Scilla, 480, 500.
 Scillipicrin, 500.
 Scillitin, 500.
 Scillitoxin, 500.
Scoparia cacumina, 470.
 Scoparin, 470.
 Sclerotic acid, 561.
 Sclerotinic acid, 561.
 Sea-salt, 88.
 Sediments, urinary, 594.
 „ urinary, microscopic examinations of, 596.
 Seidlitz powder, 875.
 Selenion, 745.
 Selenium, 852, 745.
Senega radix, 500.
Senna alexandrina, 492.
 „ *indica*, 492.
 Sepia, 571.
 Serolin, 537.
Serpentaria radix, 473.
 Serpent's excrement, 428.
 Sesamé-oil, 548.
Sesamum indicum, 548.
Sevum præparatum, 548.
 Sexivalence, 54.
 Shale, 150.
 Shark-liver oil, 548.
 Shellac, 570.
 Sherry wine, 503.
 Shot, 238.
 Sienna, 571.
 Sifting, an aid to analysis, 435.
 Silica, 418.
 Silicate of aluminium, 151.
 „ calcium, 112.
 „ lithium, 258.
 „ magnesium, 418.
Silicates, 413.
 „ quantitative estimation of, 706.
 „ tests for, 414.
 Silicio acid, 413, 415.
 „ anhydride, 413.
 Siliciuretted hydrogen, 415.
 Silico-fluoride of barium, 111.
 Silicon, chloride, 415.
 Silicon, derivation of word, 25.
 „ fluoride, 415.
 „ „ hydride, 415.
 „ „ oxide, 415.
Silver, 245.
 „ ammonio-nitrate, 198, 234.
 „ analytical reactions of, 249.
 „ antidotes to nitrate of, 250.
 „ arseniate, 197, 249.
 „ arsenite, 197.
 „ bromide, 249.
 „ chloride, 247.
 „ chromate, 250.
 „ citrate, 381.
 „ coinage, 246, 632.
 „ cyanide, 249.
 „ derivation of word, 25.
 „ estimation of, by cupellation, 694.
 „ „ extraction of, 245.
 „ „ german, 142.
 „ „ iodide, 249.
 „ „ nitrate, impure, 246.
 „ „ nitrate, pure, 247.
 „ „ oxalate, 370.
 „ „ oxide, 248.
 „ „ phosphate, 249.
 „ „ pure, 247.
 „ „ quantitative estimation of, 693.
 „ „ standard solution of nitrate of, 669.
 „ „ sulphate, 246.
 „ „ sulphide, 249.
 „ „ sulphide, native, 245.
 „ „ sulphite, 358.
 „ „ tartrate, 376.
 „ „ tree, 350.
 „ „ volumetric estimation of, 695.
 Sinalbin, 532.
 Sinapine, 533.
Sinapis, 533.
 „ „ impurities in, 741.
 „ „ *juncea*, 533.
 Singrin, 533.
 Siphon, 118.
 Sise, 540.
 Slaked lime, 114.
 Slate, 150.
 Smalt, 266, 570.
 Smilacin, 499.
 Soap, ammonium, calcium, green, hard, potassium, sodium, soft, 544.
 „ „ curd, 545.
 „ „ -stone, 572.
 „ „ -wort, 499.

- Socaloin, 531.
 Sootrine aloes, 531.
 "Soda," 83, 865.
 Soda-alum, 151.
 "ash, 91, 865.
 "caustic, 84.
 "lime, 713.
 "powders, 90.
 "solution of chlorinated, 123.
 "solution, Table of strength of, 575, 2nd edit.
 "standard solution of, 655.
 "valerianate of, 423.
 "volumetric estimation of, 651.
 "water, 88.
Soda caustica, 84.
 "caustica, impurities in, 741.
 "tartarata, 74, 89, 374.
 "tartarata, impurities in, 741.
Sodæ acetæ, 85.
 "acetæ, impurities in, 741.
 "arsenias, 188.
 "arsenias, impurities in, 741.
 "bicarbonas, 85, 366.
 "bicarbonas, impurities in, 741.
 "carbonas, 366.
 "carbonas, impurities in, 741.
 "carbonas *essiccata*, 87.
 "chloratæ liquor, 90.
 "citro-tartaræ *effervescens*, 90.
 "hypophosphis, 402.
 "hyposulphis, impurities in, 741.
 "liquor, 84.
 "nitras, 330.
 "nitras, impurities in, 741.
 "phosphas, 120.
 "phosphas, impurities in, 741.
 "sulphas, 308.
 "sulphas, impurities in, 742.
 "sulphis, 357.
 "valerianus, 423.
 "valerianus, impurities in, 742.
 Sodæ carbonate, etc., *vide* Salts of sodium.
Sodii chloridum, 83.
Sodii-citrate of iron, 168.
 "tartrate of iron, 168.
Sodium, 83.
 "acetate, 85.
 "acid carbonate, 85.
 "acid sulphate, 308.
 "analytical reactions of, 92.
 "and aluminium double chloride, 150.
 "and platinum, chloride of, 285.
 "antimoniate, 92.
 "arseniate, 188.
Sodium, arsenite, 186.
 "bicarbonate, 85.
 "bicarbonate, chemically pure, 646.
 "bisulphite, 357.
 "bromate, 91.
 "bromide, 91.
 "carbonate, 88.
 "carbonate, chemically pure, 646.
 "carbonate, manufacture of, 91, 364. [91.
 "chlorate, manufacture of, 91.
 "chloride, 83.
 "cholate, 545.
 "citrate, 91.
 "derivation of word, 24.
 "flame, 92.
 "glycocholate, 545.
 "hydrate, 84.
 "hypochlorite, 89.
 "hypophosphite, 402.
 "hyposulphite, 404.
 "in solutions of soda of different densities, Table showing the amount of oxide of, 575, 2nd edit.
 "iodate, 91.
 "iodide, 91.
 "manganate, 91.
 "nitrate, 83, 330.
 "other compounds of, 90.
 "oxalate, 369.
 "permanganate, 91.
 "phosphate, 90.
 "phosphate, how prepared from phosphate of calcium, 120.
 "potassium and ammonium, separation of, 107.
 "pyroarseniate, 188.
 "quantitative estimation of, 651, 678.
 "salts, analogy of, to potassium salts, 91.
 "salts, sources of, 83.
 "sulphate, 368.
 "sulphite, 357.
 "taurocholate, 545.
 "valerianate, 423.
 Soft soap, 544.
 Soils, analysis of, 732.
Solanin, 470.
Solanidine, 470.
Solanine, 470.
Solanum dulcamara, 470.
 "tuberosum, starch of (fig.), 476.

Solassi juice, 406.
 Solder, 238, 276.
 Solid, definition of, 57.
 Solid fats, 615.
 " potash, 68.
 Solids, to take the specific gravity of, 631 *et seq.*
 Solids lighter than water, to take the specific gravity of, 634.
 Solubility of carbonic acid gas in water, 88.
 " of gases in water, 89.
 " of precipitates in strong solutions of salts, 255.
 " or insolubility of salts in water, Table of, 430.
 Soluble cream of tartar, 390.
 " glass, 414.
 " starch, 478.
 " substances, to take the specific gravity of, 633.
 " tartar, 69.
 Solution of acetate of ammonium, 97.
 " acetate of copper, 216.
 " acetate of potassium, 70.
 " acetate of sodium, 85.
 " albumen, 535.
 " ammonia, 96.
 " ammonio-nitrate of silver, 198.
 " ammonio-sulphate of copper, 198.
 " ammonio-sulphate of magnesium, 667.
 " arsenic in acid, 186.
 " arsenic in alkali, 185.
 " boric acid, 390.
 " bromine, 313.
 " carbonate of ammonium, 99.
 " chloride of ammonium, 95.
 " chloride of antimony, 201.
 " chloride of barium, 109.
 " chloride of calcium, 113.
 " chloride of calcium, saturated, 113.
 " chloride of gold, 261.
 " chloride of tin, 277.
 " chloride of zinc, 145.
 " chlorinated lime, 123.
 " chlorinated soda, 90.
 " chlorine, 19, 308.

Solution of citrate of ammonium, 97.
 " ferricyanide of potassium, 400.
 " ferrocyanide of potassium, 398.
 " gelatine, 540.
 " iodate of potassium, 76, 344.
 " iodide of potassium, 76.
 " iodine, 316.
 " lime, 115.
 " litmus, 101.
 " nitrate of mercury, 235.
 " oxalate of ammonium, 100.
 " oxyacetate of lead, 340.
 " perchloride of iron, 163, 164.
 " perchloride of mercury, 237.
 " perchloride of platinum, 263.
 " permnitrate of iron, 173.
 " persulphate of iron, 164.
 " phosphate of sodium, 131.
 " phosphoric acid, 337.
 " potash, 63.
 " red prussiate of potash, 400.
 " soda, 84.
 " strychnia, 461.
 " sulphate of calcium, 134.
 " sulphate of indigo, 337.
 " sulphate of iron, 157.
 " sulphide of ammonium, 101.
 " sulphhydrate of ammonium, 101.
 " tartaric acid, 373.
 " yellow prussiate of potash, 398.
 Sonnenschein's process for poisonous alkaloids, 587.
 Soot, 23.
 Soubresauts, 324.
 Source of heat, 6.
 Sovereign, weight of the, 230.
Soymida febrifuga, 473.
 Spanish liquorice, 406.
 Spar, fluor-, 113.
 " heavy, 110.

- Sparteia or sparteine, 470.
 Spathic iron-ore, 154.
 Spearmint-oil, 555.
 Specific gravity, 618, 627.
 " gravity bottles, 629.
 " gravity of gases, 634.
 " gravity of liquids, 632.
 " gravity of official liquids, 629.
 " gravity of oxygen, 14.
 " gravity of powders, 633.
 " gravity of solids, 632.
 " gravity of solids lighter than water, 634.
 " gravity of soluble substances, 633.
 " heat, 140, 638.
 " weight, 637.
 Spectroscope, 301, 576.
 Spectrum analysis, 301, 576.
 Specular iron-ore, 154.
 Speculum metal, 276.
 Speiss, 268.
 Spermaceti, 546.
 Spermatozoa in urine, 601.
 Sperm-oil, 546.
Spiraea ulmaria, 499, 526.
 Spirit, methylated, 517.
 " of French wine, 507.
 " of myrcia, 503.
 " of nitrous ether, 511.
 " of nitrous ether, adulterated, 518.
 " of turpentine, 557.
 " of wine, 504.
 " of wine, impurities in, 742.
 " proof, 504.
 " pyroxylic, 517.
 " rectified, 139, 504.
 " wood-, 517.
 Spirits, 605.
 " analysis of, 728.
Spiritus ætheris, 511.
 " *ætheris nitrosi*, 410, 511.
 " *ætheris nitrosi*, impurities in, 742.
 " *ammoniac aromaticus*, 98.
 " *ammoniac aromaticus*, impurities in, 742.
 " *ammoniac fatidus*, 99.
 " *armoriacæ compositus*, 553.
 " *cajuputi*, 532.
 " *chloroformi*, impurities in, 742.
 " *juniiperi*, 552.
 " *lavandulæ*, 552.
 " *mentha piperitis*, 552.
 " *myrcia*, 503.
Spiritus myristica, 552.
 " *rectificatus*, 504.
 " *rectificatus*, impurities in, 742.
 " *rosmarini*, 552.
 " *tenuior*, 504.
 " *tenuior*, impurities in, 742.
 " *vini gallici*, 507.
 Spodumene, 258.
 Spogel seeds, 480.
 Spongy platinum, 285.
 Spontaneous combustion, 175.
 Spotted cranesbill, 420.
 Spruce fir, 562.
 Spurge laurel, 562.
Squalus carcharias, 548.
 Squill, 480, 500.
 Standard gold, 28.
 Standard solution of hyposulphite of sodium, 670.
 " solution of iodine, 663.
 " solution of nitrate of silver, 659.
 " solution of oxalic acid, 646.
 " solution of red chromate of potassium, 665.
 " solution of soda, 655.
 " solution of sulphuric acid, 654.
 Stannate of sodium, 277.
 Stannates, 278.
 Stannic acid, 278.
 " anhydride, 277.
 " chloride, 277.
 " oxide, 277.
 " sulphide, 279.
 " sulphide, anhydrous, 279.
 Stannous chloride, 277.
 " chloride, solid, 277.
 " hydrate, 279.
 " oxide, 279.
 " sulphide, 278.
 Stannum, 26.
 Star-anise oil, 553.
 Starch, 473.
 " action of diastase upon, 473.
 " action of dilute acids upon, 473.
 " blue, 474.
 " granules, composition of, 475.
 " iodide of, 475.
 " potato, 473.
 " quantitative estimation of, 726.
 " soluble, 478.
 " white, 473.

- Starches, microscopy of, 475.
 Stas's process for poisonous alkaloïds, 586.
 Stavesacre, 467.
 Steam-bath, 120.
 Stearic acid, 544, 549.
 Stearine, 544.
 Stearoptens, 550.
 Steatite, 572.
 Steel, 154.
 " wine, 172.
 Stibium, 25.
 Stick liquorice, 496.
 Still, 137.
 Stone-coal, 275.
 " red, 569.
 Storax, 566.
 Stout, 508.
Stramonii folia, 468.
 " *semina*, 468.
 Strasburg turpentine, 557.
 Strawberry, sugar in, 496.
 Stream-tin, 277.
 Strontianite, 259.
 Strontium, 259.
 " analytical reactions of, 260.
 " carbonate, 259.
 " derivation of word, 25.
 " flame, 261.
 " nitrate, 260.
 " sulphate, 260.
 Structure of flame, 13.
 Strychnia or strychnine, 459.
 " or strychnaine, impurities in, 742.
 " analytical reactions of, 460.
 " in organic mixtures, detection of, 588.
Strychnos ignatiua, 459.
 " *nux vomica*, 459.
 Styraçin, 566.
Styrax benzoin, 566.
 " *preparatus*, 566.
 Styrol, 566.
 Styronc, 566.
 Subacetate of copper, 216.
 " of lead, 239.
 Subchloride of mercury, 228.
 Sublimation, 97.
 Sublimed sulphur, 251.
 Subnitrate of bismuth, 288.
 Substances readily deoxidized, quantitative estimation of, 669.
 Substances readily oxidized, quantitative estimation of, 662.
 Substitution products, 520.
Succi, 606.
 Succinate of potassium, 416.
 Succinic acid, 415.
Succinum, 415.
Succus limonium, 379.
 Sucrose, 486.
 Suet, 546.
 " prepared, 546.
 Sugar, 484.
 " action of alkali upon, 487.
 " amount in various fruits, 486.
 " brown, 485.
 " candy, 485.
 " -cane, 484, 485.
 " detection of in urine, 591.
 " grape-, 484.
 " from starch, 487.
 " inverted, 484.
 " lump, 485.
 " -maple, 485.
 " milk-, 486.
 " moist, 485.
 " of gelatine, 545.
 " of lead, 239.
 " patent, 488.
 " quantitative estimation of, 725.
 " test for, 484.
 Sulphate of aluminium, 151.
 " aluminium and ammonium, 150.
 " ammonium, 95.
 " barium, 110.
 " bismuth, 289.
 " calcium, 102, 119.
 " chromium, 262.
 " cobalt, 267.
 " copper, 215.
 " copper, anhydrous, 216.
 " cupr-diammon-diammonium, 234.
 " indigo, 237.
 " iron, 156.
 " iron and ammonium, 151.
 " iron, solution of, 157.
 " lead, 244.
 " lithium, 259.
 " magnesium, 126.
 " mercury, 228.
 " potassium, 74.
 " quinine, 453.
 " sodium, 206.
 " strontium, 200.
 " zinc, 142.
 Sulphates, 358.
 " analytical reactions of, 362.

Sulphates, quantitative estimation of, 701.

Sulphethylic acid, 510.

Sulphide of allyl, 501.

- " ammonium, 101.
- " antimony, 200, 206.
- " arsenicum, 185, 196.
- " arsenicum, native, 185.
- " barium, 110.
- " bismuth, 291.
- " cadmium, 286.
- " cobalt, 267.
- " iron, 23, 160, 177, 178.
- " lead, 248.
- " lead, native, 237.
- " manganese, 265.
- " mercury, 234.
- " mercury, native, 218.
- " nickel, 263.
- " potassium, 68.
- " platinum, 283.
- " silver, 249.
- " silver, native, 245.
- " zinc, 147.
- " zinc, native, 142.

Sulphides, 850.

- " analytical reactions of, 354.
- " quantitative estimation of, 700.

Sulphindigotic acid, 337.

Sulphindyllic acid, 337.

Sulphite of barium, 358.

- " calcium, 357.
- " silver, 353.
- " sodium, 357.
- " zinc, 147.

Sulphites, 355.

- " analytical reactions of, 357.
- " quantitative estimation of, 700.

Sulphocarbonates, 530.

Sulphocarbonates, 366.

Sulphocarbolic acid, 530.

Sulphocyanate of acrinyl, 532.

- " allyl, 532, 556.
- " butyl, 516, 553.
- " iron, 179, 416.
- " mercury, 417.

Sulphocyanates, 416.

Sulphocyanic acid, 416.

Sulphocyanides, 417.

Sulphocyanogen, 417.

Sulphophenates, 530.

Sulphophenic acid, 530.

Sulphosalicylic acid, 531.

Sulphostannates, 279.

Sulphovinic acid, 361, 510.

Sulphur, 21, 350.

- " adulteration of, 353.
- " allotropy of, 351, 432.
- " analogies, 336.
- " analytical reactions of, 354.
- " arsenic in, 196.
- " bromide, 354.
- " chloride, 354.
- " derivation of word, 23.
- " estimation of, 700.
- " flowers of, 351.
- " hypochloride, 355.
- " iodide, 316.
- " liver of, 68.
- " milk of, 352.
- " oxyacids, 404.
- " plastic, 351.
- " precipitated, 352.
- " roll, 351.
- " sublimed, 351.

Sulphur lotum, 351.

- " *precipitatum*, 352.
- " *precipitatum*, impurities in, 742.
- " *sublimatum*, 351.
- " *sublimatum*, impurities in, 742.

Sulphurated antimony, 204.

- " potash, 68.

Sulphurets, *vide* Sulphides.

Sulphuretted hydrogen, 102, 350.

Sulphuric acid, 358.

- " acid, antidotes to, 363.
- " acid, aromatic, 361.
- " acid, estimation of, in vinegar, 702.
- " acid, dilute, 261.
- " acid, fuming, 362.
- " acid, Nordhausen, 362.
- " acid in organic mixtures, detection of, 578, 581.
- " acid, purification of, 361.
- " acid, standard solution of, 654.
- " acid and sulphuric anhydride in dilute acid of different densities, Tables showing percent. of, 566, 2nd edit.
- " acid, volumetric estimation of, 657.
- " anhydride, 361.

Sulphuris iodidum, 316.

- " *iodidum*, impurities in, 742.

Sulphurous acid, 21, 355.

- Sulphurous acid, volumetric estimation of, 663.
 anhydride, 355.
 Sulphydrate of ammonium, solution of, 101.
 Sulphydic acid, 350.
 Sumatra camphor, 558.
 Sumbul, 564.
 root, 564.
 Sumbul radice, 564.
 Superphosphate of lime, 383.
 Supporters of combustion, 12.
Suppositoria acidi tannici, 418.
 " *morphice*, 449.
 " *plumbi composita*, 229.
 Suppositories, 606.
 Surface unit, 621.
 Surgery, 2.
 Sweetbread, 542.
 Sweet-flag oil, 556.
 " spirit of nitre, 410, 511.
 " spirit of nitre, adulterated, 518.
 Sylvic acid, 560.
 Symbol, function of, 56.
 Symbols of elements, 23 *et seq.*
 " illustration of chemical action by, 42.
 Sympathetic inks, 268.
 Synaptase, 491.
 Synthesis, 62, 114.
 Syphon, *see* Siphon.
 Syrup of iodide of iron, 22.
Syrupi, impurities in, 742.
 Syrups, 606.
 " specific gravities of, 630.
Syrupus aurantii, 553.
 " *aurantii floris*, 553.
 " *ferri iodidi*, 22.
 " *ferri phosphatis*, 159.
Tabaci folia, 469.
 Tables, various, *see* Appendix.
 Talc, 150.
Tamarindus, 378.
 " impurities in, 742.
 Tannic acid or tannin, 417.
 Tanning, 418.
 Tantalum, 745.
 Tapioca, 474.
 " starch (fig.), 476.
Taraxaci radice, 473.
 Taraxacin, 473.
 Tartar, cream of, 57, 61, 73, 372.
 " emetic, 203.
 " emetic, estimation of anti-mony in, 688.
 " meaning of, 372.
 Tartarated antimony, 203.
 Tartaric acid, 371, 373.
 " saturating power of, 374, 743.
 " solution of, 373.
Tartarus boracatus, 391.
 Tartrate of ammonium, 105.
 " antimony and potassium, 303, 372.
 " calcium, 375.
 " potassium, acid, 61, 73, 81, 373.
 " potassium, neutral, 74, 374.
 " potassium and sodium, 89, 374.
 " silver, 376.
 " sodium, 375.
 Tartrates, 372.
 " analytical reactions of, 375.
 " volumetric estimation of, 653.
 Taurine, 545.
 Taurocholates, 545.
 Tea, 470.
 Teal oil, 548.
 Telini fly, 558.
 Tellurium, 745.
 Temperature, correction of vol. of gas for, 635.
 " quantitative determination of, 611.
 Terbium, 745.
Terebinthina canadensis, 557.
 Terpenes, 550.
 Terra di sienna, 569.
Terra japonica, 419.
 Test-papers, 101.
 " -tube, 5.
 Tetrads, 133.
 Tetramines, 447.
 Tetrathionic acid, 406.
 Tetryl, 515.
 Thalleioquin, 456.
 Thallium, 745.
 Thebaia, 450.
 Theia, 470.
 Theine, 470.
 Thenard's-blue, 570.
 Theobroma-oil, 546.
 Theobromate of glyceryl, 546.
 Therapeutics, definition and derivation of, 3.
Thoriaca, 489.
 Thermolysis, 638.
 Thermometers, 612.
 " Celsius's, 612.

- Thermometers, Centigrade, 613.
 " Fahrenheit's, 613.
 " Réaumur's, 613.
 Thermometric scales, conversion of
 degrees of, 614.
 Thionic acids, 404.
 Thiosulphates, 404.
 Thorium, 745.
 Thorium, 745.
 Thorn-apple, 468.
Thus americanum, 564.
 masculum, 565.
 Thyme, 557.
 Thymene, 557.
 Thymol, 553, 557.
Thymus vulgaris, 555, 557.
 Tiglic acid, 548.
 Tin, 275.
 " amalgam, 276.
 " analytical reactions of, 278.
 " antidotes to, 280.
 " block, 276.
 " chloride, 277.
 " derivation of word, 26.
 " dropped or grain, 276.
 " foil, 276.
 " granulated, 276.
 " oxide, 277.
 " perchloride, 277.
 " plate, 276.
 " prepare-liquor, 278.
 " -stone, 275.
 " tacks, 276.
 " -white cobalt, 266.
 Tincal, 382.
Tinctura ferri acetatis, 165.
 " *ferri perchloridi*, 163.
 " *iodi*, 316.
 " *quinia*, 455.
 " *quinia ammoniata*, 455.
Tinctura, 163.
 Tinctures, 163, 605.
 Tinnevely senna, 492.
Tinospora cordifolia, 473.
 Titanium, 745.
 Tobacco, 469.
Toddalia radis, 473.
 Toluene, 566.
 Tolu balsam of, 566.
 Toluol, 528.
 Toxicology, 577.
 Tragacanth, 123.
 Treacle, 489.
 Tree, lead, 244.
 " silver, 250.
 Trade, 133.
 Triamines, 447.
 Triangle, wire, 105.
 Tribasic acids, 304.
 Tribasylous radicals, 304.
 Triethylamine, 447.
 Triethylia, 447.
 Trimethylamine, 447.
 Trinitro-carbolic acid, 530.
 Trinitrocellulin, 482.
 Triphane, 258.
 Tripoli, 413.
 Trithionic acid, 405.
Triticum, starch of (fig.), 476.
 Trityl, 515.
 Tritylia, 447.
 Trivalence, 54.
 Trivalent radicals, 54, 66, 133.
Trochisci acidi tannici, 418.
 " *bismuthi*, 289.
 " *ferri redacti*, 175.
 " *morphice*, 449.
 " *morphice et ipecacuanha*,
 449.
 " *potassæ chloratis*, 342.
 sodæ bicarbonatis, 89.
 Tropate of atropine, 464.
 Tropic acid, 464.
 Tropidine, 464.
 Tropine, 464.
 Tube-funnels, 10.
 Tubes for collecting gases, 6.
 " glass, see Glass tubes.
 Tungsten, 745.
 Turgite, 167.
 Turkey corn, 467.
 Turmeric, 168.
 " oil, 557.
 " paper, 101.
 Turnbull's blue, 177, 400.
 Turnsole, 570.
 Turpentine, 556, 564.
 " American, 557, 564.
 " Bordeaux, 557.
 " Canadian, 557.
 " Chian, 557.
 " French, 557.
 " rectified oil of, 557.
 " spirit of, 557.
 " Strasburg, 557.
 " Venice, 557.
 Turpeth mineral, 226.
 Turps, 557.
Tylophora asthmatica, 468.
 Type-metal, 200, 238, 287.
 Types, chemical, 443.
 Typical formulæ, 443.

Ulm cortex, 419, 480.
 " *fulvæ cortex*, 480.
 Ultimate analysis, 709.

- Ultramarine blue, 570.
 " green, 571.
 Umbelliferone, 565.
 UMBER, 571.
Uncaria gambier, 419.
Unguentum aconitiae, 463.
 " *antimonii tartarati*, 203.
 " *atropiae*, 464.
 " *cadmii iodidi*, 236.
 " *cerussae*, 238.
 " *hydrargyri*, 219.
 " *hydrargyri ammoniati*, 233.
 " *hydrargyri iodidi rubri*, 234.
 " *hydrargyri nitratis*, 225.
 " *hydrargyri oxidi rubri*, 230.
 " *hydrargyri subchloridi*, 229.
 " *indi*, 316.
 " *plumbi acetatis*, 239.
 " *plumbi carbonatis*, 238.
 " *plumbi iodidi*, 241.
 " *plumbi subacetatis compositum*, 240.
 " *potassae sulphuratae*, 68.
 " *sulphuris iodidi*, 316.
 " *veratrinae*, 472.
 " *sinct*, 146.
 Units of length, surface, capacity and weight, 621.
 Univalence, 54.
 Univalent radicals, 54, 133.
 Uralium, 745.
 Uranium, 745.
 Urari, 461.
 Urate of lithium, 259.
 Urates, 422.
Urceola elastica, 567.
 Urea, 395, 491, 593.
 " artificial, 395, 593.
 " nitrate of, 592.
 " test for excess of in urines, 592.
 Uric acid, 422, 595.
 Urinary calculi, 601.
 " calculi, examination of, 601.
 " deposits or sediments, plates of, 597, 599.
 " sediments, 594.
 " sediments, microscopical examination of, 596.
 Urine, 422, 590.
 " diabetic, 592.
 " estimation of sugar in, 726.
 " estimation of urea in, 593.
 " morbid, examination of, 591.
 Urinometer, 592.
- Uvae*, 372, 488.
 " *ursi folia*, 420.
 Valerene, 557.
 Valerian oil, 557.
Valeriana radia, 557.
 Valerianate of amyl, 424, 526.
 " sodium, 423.
 " zinc, 147, 424.
 Valerianates, 423.
 Valerianic acid, 423, 549, 557.
 Valerol, 557, 564.
 Vanadates, 389.
 Vanadinite, 389.
 Vanadium, 389, 745.
 " relationship to nitrogen, phosphorus and arsenicum, 389.
 Vanilla, 425.
Vanilla planifolia, 425.
 Vanillic acid, 425.
 Vanillin, 425.
Vapor acidi hydrocyanici, 326.
 " *chlori*, 20.
 " *conice*, 467.
 " *iodi*, 315.
 Vapour-density, 635.
Variolaria, 570.
 Vegetable albumen, 539.
 " and animal life, relation of, 9.
 " casein, 474, 539.
 " crocus, 568.
 " fibrin, 474, 539.
 " gelatine, 479.
 " green, 571.
 " jelly, 479.
 " oil, 544.
 " rouge, 570.
 " substances, 445 *et seq.*
 Venetian red, 168.
 Venice turpentine, 557.
Veratralbia, 468.
Veratri viridis radia, 468.
Veratrina, 471.
 " impurities in, 742.
 Veratria or Veratrine, 468, 471, 742.
Veratrum album, 468.
 " *viride*, 468.
 Verbena oil, 557.
 Verdigris, 216.
 Vermilion, 234, 569.
Veronica virginica, 473.
 Vinegar, 346.
 " estimation of mineral acids in, 702.
 " impurities in, 735.
 " of cantharides, 346.

Vinegar of squill, 346.

Vinum antimoniale, 203.

„ *aurantii*, 503.

„ *ferri*, 172.

„ *ferri citratis*, 172.

„ *quinia*, 455.

„ *sericum*, 503.

Vitriol, blue, 157, 215.

„ green, 157, 360.

„ oil of, 360, 361.

„ white, 157.

Volatile oils, *see* Oils.

Volatility of salts of ammonium, 105.

Volatilization, 105.

Volcanic ammonia, 95.

Volume, combination by, 50.

„ of gas, corrections of, 635.

„ molecular, 533.

Volumetric quantitative analysis,
609.

„ estimation of acetate
of lead, 660.

„ estimation of acetic
acid, 656.

„ estimation of acids,
655.

„ estimation of alkalies,
646.

„ estimation of alkaline
carbonates, 649, 651.

„ estimation of ammo-
nia solutions, 647.

„ estimation of arseni-
ate of iron, 667.

„ estimation of arseni-
ate of sodium, 661.

„ estimation of arsenic
and arsenical solu-
tions, 665.

„ estimation of borax,
649.

„ estimation of bro-
mide of potassium,
661.

„ estimation of bro-
mides, 661.

„ estimation of chlo-
rides, 695.

„ estimation of chlori-
nated lime, 672.

„ estimation of chlori-
nated soda, 672.

„ estimation of chlo-
rine, 659.

„ estimation of citrate
of potassium, 652.

„ estimation of citrates,
652.

**Volumetric estimation of cyan-
ides, 659.**

„ estimation of hydro-
chloric acid, 657.

„ estimation of hydro-
cyanic acid, 659.

„ estimation of hypo-
sulphite of sodium,
665.

„ estimation of mag-
netic oxide of iron,
668.

„ estimation of nitric
acid, 657.

„ estimation of official
compounds, 647 *et*
seq.

„ estimation of phos-
phate of iron, 667.

„ estimation of potash,
651.

„ estimation of saocha-
rated carbonate of
iron, 668.

„ estimation of silver,
656.

„ estimation of soda,
651.

„ estimation of sugar,
725.

„ estimation of sul-
phides, 700.

„ estimation of sul-
phites, 664.

„ estimation of sulphu-
ric acid, 657.

„ estimation of sulphu-
rous acid, 663.

„ estimation of tar-
trates, 652.

„ solutions, 646, 654,
655, 658, 663, 665,
670.

Vulcanite, 567.

Vulcanized india-rubber, 567.

Wahoo bark, 473.

Warmth of animals, how kept up, 9.

Wasp sting, 395.

Washing-bottles, 117.

„ precipitates, 117.

Water, aerated, 88.

„ aspirator, 360.

„ ammonia in potable, 648.

„ bath, 116, 119.

„ boiling-point of, 615.

„ chalybeate, 158.

„ composition of, 12.

- Water, cubic inches of, in a gallon, 636.
 „ estimation of, 708.
 „ evaporation of, 71.
 „ formation of, expressed by symbols, 42.
 „ hardness of, 367.
 „ hemlock, 555.
 „ lead in, 243.
 „ lime, 115.
 „ nitrites in, 409.
 „ of crystallization, 87.
 „ of crystallization, quantitative estimation of, 708.
 „ -oven, 675.
 „ oxygenated, 110.
 „ preparation of, 11.
 „ purification of, 137, 367.
 „ weight of a cubic inch of, 636.
 „ weight of minim, drachm, ounce, pint, and gallon, 626.
 „ softness of, 367.
 „ type, 443.
 Wax, 546.
 Weighing-tubes, 675.
 Weight, 618.
 „ estimation of, 618.
 „ molecular, 53.
 „ of air, 637.
 „ of hydrogen, 637.
 „ of water, 637.
 „ specific, 627.
 Weights, atomic, 49.
 „ balance, 619.
 „ and measures, 619.
 „ and measures of the metric decimal system, 620*et seq.*
 „ and measures of the British Pharmacopœia, of 1867, 626.
 „ relative, 49.
 Weld, 569.
 Welding, 154, 285.
 Wheaten flour, 474.
 Wheat-starch (fig.), 476.
 Whey, 486, 537.
 Whisky, 503.
 White arsenic, 177.
 „ indigo, 337.
 „ lead, 238, 693.
 „ pepper, 469.
 „ pigments, 571.
 „ precipitate, 232.
 „ precipitate, fusible, 233.
 „ precipitate, infusible, 233.
 „ resin, 560.
 White vitriol, 157.
 „ wax, 546.
 Whiting, 118.
 Whortleberry, sugar in, 486.
 Wild indigo, 473.
 Willow-bark, 406, 498.
 Wine, 503, 687.
 „ antimonial, 203.
 „ iron, 172.
 „ orange, 503.
 „ quinine, 455.
 „ sherry, 503.
 „ steel, 172.
 Winter-green, oil of, 526.
 Wire-gauze tray, 20.
 „ triangle, 106.
 Witherite, 110.
 Wood-charcoal, 120.
 „ -creasote, 529.
 „ -naphtha, 517.
 „ -oil, 563.
 „ -spirit, 517.
 „ -tar, 564.
 Woody night-shade, 470.
 Wormwood, 554.
 Woorara, 461.
 Wourali, 461.
 Wrought iron, 154.
Xanthorrhiza apiifolia, 465.
Xanthoxylon frazineum, 465.
 Xanthin, 603.
 Xylol, 528.
 Yard, 626.
 Yeast, 501, 502.
 Yolk of egg, 536.
 Yellow chromate of potassium, 111.
 „ colouring-matters, 563.
 „ dock, 395.
 „ ochre, 568.
 „ oxide of mercury, 230.
 „ parilla, 465.
 „ prussiate of potassium, 323, 397.
 „ root, 465.
 „ sienna, 563.
 „ wax, 546.
 „ wood, 563.
 Yolk of egg, 536.
 Ytterbium, 745.
 Yttrium, 745.
 Zanaloin, 531.
 Zaffre, 266.
 Zanzibar aloes, 531.
 Zea, starch of (fig.), 476.
 Zinc, 124.

- Zinc, acetate, 145.**
 „ analytical reactions of, 147.
 „ antidotes to, 148.
 „ carbonate, 142, 145.
 „ chloride, 143.
 „ derivation of word, 24.
 „ detection of, in presence of aluminium and iron, 180.
 „ -ethyl, 514.
 „ ferrocyanide of, 148.
 „ granulated, 10.
 „ hydrate, 147.
 „ in organic mixtures, detection of, 579.
 „ oxide, 146.
 „ oxide, Hubuck's, 146.
 „ quantitative estimation of, 684.
 „ sulphate, 142.
 „ sulphide, 147.
 „ sulphide, native, 142.
- Zinc, sulphite, 147.**
 „ valerianate, 147, 424.
 „ white, 145.
Zinci acetas, 146.
 „ acetas, impurities in, 742.
 „ carbonas, 145.
 „ carbonas, impurities in, 742.
 „ chloridi liquor, 145.
 „ chloridum, 144.
 „ chloridum, impurities in, 742.
 „ oxidum, 146.
 „ oxidum, impurities in, 742.
 „ sulphas, 142.
 „ sulphas, impurities in, 742.
 „ unguentum, 146.
 „ valerianas, 147, 424.
 „ valerianas, impurities in, 742.
Zincum, 142.
 „ granulatum, 10.
Zingiber, 557.
Zirconium, 745.

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